

3. Energy

Energy-related activities were the primary sources of U.S. anthropogenic greenhouse gas emissions, accounting for 86 percent of total emissions on a carbon equivalent basis in 2003. This included 97, 39, and 15 percent of the nation's carbon dioxide (CO₂), methane (CH₄), and nitrous oxide (N₂O) emissions, respectively. Energy-related CO₂ emissions alone constituted 83 percent of national emissions from all sources on a carbon equivalent basis, while the non-CO₂ emissions from energy-related activities represented a much smaller portion of total national emissions (4 percent collectively).

Emissions from fossil fuel combustion comprise the vast majority of energy-related emissions, with CO₂ being the primary gas emitted (see Figure 3-1). Globally, approximately 24,240 Tg CO₂ were added to the atmosphere through the combustion of fossil fuels in 2000, of which the United States accounted for about 23 percent.¹ Due to the relative importance of fossil fuel combustion-related CO₂ emissions, they are considered separately, and in more detail than other energy-related emissions (see Figure 3-2). Fossil fuel combustion also emits CH₄ and N₂O, as well as ambient air pollutants such as nitrogen oxides (NO_x), carbon monoxide (CO), and non-methane volatile organic compounds (NMVOCs). Mobile fossil fuel combustion was the second largest source of N₂O emissions in the United States, and overall energy-related activities were collectively the largest source of these ambient air pollutant emissions.

Figure 3-1: 2003 Energy Sector Greenhouse Gas Sources

Figure 3-2: 2003 U.S. Fossil Carbon Flows (Tg CO₂ Eq.)

Energy-related activities other than fuel combustion, such as the production, transmission, storage, and distribution of fossil fuels, also emit greenhouse gases. These emissions consist primarily of fugitive CH₄ from natural gas systems, petroleum systems, and coal mining. Smaller quantities of CO₂, CO, NMVOCs, and NO_x are also emitted.

The combustion of biomass and biomass-based fuels also emits greenhouse gases. Carbon dioxide emissions from these activities, however, are not included in national emissions totals because biomass fuels are of biogenic origin. It is assumed that the carbon released during the consumption of biomass is recycled as U.S. forests and crops regenerate, causing no net addition of CO₂ to the atmosphere. The net impacts of land-use and forestry activities on the carbon cycle are accounted for within the Land-Use Change and Forestry sector. Emissions of other greenhouse gases from the combustion of biomass and biomass-based fuels are included in national totals under stationary and mobile combustion.

Table 3-1 summarizes emissions from the Energy sector in units of teragrams of CO₂ equivalents (Tg CO₂ Eq.), while unweighted gas emissions in gigagrams (Gg) are provided in Table 3-2. Overall, emissions due to energy-related activities were 5,963.4 Tg CO₂ Eq. in 2003, an increase of 16 percent since 1990.

Table 3-1: Emissions from Energy (Tg CO₂ Eq.)

Gas/Source	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	4,836.4	5,409.1	5,437.7	5,512.1	5,693.5	5,592.9	5,645.3	5,694.3
Fossil Fuel Combustion	4,711.7	5,263.2	5,278.7	5,345.9	5,545.1	5,448.0	5,501.4	5,551.6
Non-Energy Use of Fuels	108.0	120.3	135.4	141.6	124.7	120.1	118.8	118.0
Waste Combustion	10.9	17.8	17.1	17.6	18.0	18.8	18.8	18.8

¹ Global CO₂ emissions from fossil fuel combustion were taken from Marland *et al.* (2003) <http://cdiac.esd.ornl.gov/trends/emis/meth_reg.htm>.

Natural Gas Flaring	5.8	7.9	6.6	6.9	5.8	6.1	6.2	6.0
<i>Biomass-Wood*</i>	212.5	226.3	209.5	214.3	217.6	190.8	195.8	201.0
<i>International Bunker Fuels*</i>	113.5	109.9	114.6	105.3	101.4	97.9	89.5	84.2
<i>Biomass-Ethanol*</i>	4.2	7.0	7.7	8.0	9.2	9.7	11.5	15.8
CH₄	248.9	234.6	230.9	222.1	224.3	221.6	215.8	212.7
Natural Gas Systems	128.3	133.6	131.8	127.4	132.1	131.8	130.6	125.9
Coal Mining	81.9	62.6	62.8	58.9	56.2	55.6	52.4	53.8
Petroleum Systems	20.0	18.8	18.5	17.8	17.6	17.4	17.1	17.1
Stationary Sources	7.8	7.4	6.9	7.1	7.3	6.7	6.4	6.7
Mobile Sources	4.8	4.0	3.9	3.6	3.4	3.1	2.9	2.7
Abandoned Coal Mines	6.1	8.1	7.2	7.3	7.7	6.9	6.4	6.4
<i>International Bunker Fuels*</i>	0.2	0.1	0.2	0.1	0.1	0.1	0.1	0.1
N₂O	56.4	69.1	69.1	68.4	67.5	62.8	59.6	56.4
Mobile Sources	43.7	55.2	55.3	54.6	53.2	49.0	45.6	42.1
Stationary Sources	12.3	13.5	13.4	13.5	14.0	13.5	13.5	13.8
Waste Combustion	0.4	0.4	0.3	0.3	0.4	0.4	0.5	0.5
<i>International Bunker Fuels*</i>	1.0	1.0	1.0	0.9	0.9	0.9	0.8	0.8
Total	5,141.7	5,712.8	5,737.7	5,802.6	5,985.3	5,877.3	5,920.7	5,963.4

* These values are presented for informational purposes only and are not included or are already accounted for in totals.

Note: Totals may not sum due to independent rounding.

Table 3-2: Emissions from Energy (Gg)

Gas/Source	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	4,836,430	5,409,100	5,437,734	5,512,062	5,693,544	5,592,947	5,645,251	5,694,332
Fossil Fuel Combustion	4,711,741	5,263,164	5,278,721	5,345,904	5,545,083	5,447,969	5,501,427	5,551,580
Non-Energy Use of Fuels	107,964.9	120,300.6	135,352.1	141,582.9	124,713.8	120,104.1	118,810.6	118,001.4
Waste Combustion	10,919	17,761	17,094	17,632	17,979	18,781	18,781	18,781
Natural Gas Flaring	5,805	7,874	6,566	6,943	5,769	6,094	6,233	5,970
<i>Biomass-Wood*</i>	212,547	226,265	209,490	214,323	217,577	190,776	195,776	201,042
<i>International Bunker Fuels*</i>	113,503	109,858	114,557	105,294	101,404	97,865	89,489	84,193
<i>Biomass-Ethanol*</i>	4,155	6,978	7,711	8,017	9,188	9,701	11,473	15,771
CH₄	11,852	11,170	10,997	10,577	10,680	10,551	10,279	10,126
Natural Gas Systems	6,112	6,363	6,276	6,066	6,289	6,277	6,221	5,998
Coal Mining	3,900	2,983	2,989	2,805	2,677	2,647	2,497	2,561
Petroleum Systems	951	895	879	848	836	831	815	815
Stationary Sources	373	351	328	338	349	318	305	319
Mobile Sources	228	193	185	172	161	147	138	128
Abandoned Coal Mines	288	385	341	349	369	331	303	306
<i>International Bunker Fuels*</i>	8	7	7	6	6	5	4	4
N₂O	182	223	223	221	218	203	192	182
Mobile Combustion	141	178	179	176	171	158	147	136
Stationary Combustion	40	44	43	43	45	43	44	45
Waste Combustion	1	1	1	1	1	1	1	1
<i>International Bunker Fuels*</i>	3	3	3	3	3	3	3	2

* These values are presented for informational purposes only and are not included or are already accounted for in totals.

Note: Totals may not sum due to independent rounding.

3.1. Carbon Dioxide Emissions from Fossil Fuel Combustion (IPCC Source Category 1A)

Carbon dioxide emissions from fossil fuel combustion in 2003 increased 0.9 percent from the previous year. This increase is primarily a result of increased demand for fuels due to a growing economy, fuel-switching from natural gas to coal in the electric power sector, and increased use of heating fuels in the residential sector caused by a colder

winter. In 2003, CO₂ emissions from fossil fuel combustion were 5,551.6 Tg CO₂ Eq., or 18 percent above emissions in 1990 (see Table 3-3).²

Table 3-3: CO₂ Emissions from Fossil Fuel Combustion by Fuel Type and Sector (Tg CO₂ Eq.)

Fuel/Sector	1990	1997	1998	1999	2000	2001	2002	2003
Coal	1,680.9	1,920.2	1,936.5	1,938.2	2,025.6	1,959.2	1,973.1	2,013.8
Residential	2.4	1.5	1.2	1.3	1.1	1.1	1.0	1.1
Commercial	12.1	12.2	8.7	9.7	8.6	9.2	8.6	9.3
Industrial	152.6	147.2	139.2	133.8	135.0	130.6	122.7	123.4
Transportation	NE	NE	NE	NE	NE	NE	NE	NE
Electricity Generation	1,513.0	1,758.4	1,786.4	1,792.4	1,880.0	1,817.4	1,839.7	1,876.3
U.S. Territories	0.6	1.0	1.0	0.9	0.9	0.9	1.2	3.6
Natural Gas	1,009.5	1,193.7	1,169.9	1,173.6	1,224.1	1,174.4	1,214.0	1,170.3
Residential	238.8	270.2	246.5	256.5	270.3	259.7	265.9	277.3
Commercial	142.6	174.3	163.5	165.2	172.4	164.5	168.7	170.7
Industrial	416.3	489.1	476.7	456.4	464.6	426.0	435.5	407.9
Transportation	35.9	41.1	35.1	35.6	35.5	33.9	37.1	35.4
Electricity Generation	176.0	218.9	248.0	259.9	280.7	289.1	305.6	277.6
U.S. Territories	NO	NO	NO	NO	0.7	1.2	1.2	1.4
Petroleum	2,020.9	2,148.9	2,172.0	2,233.7	2,295.0	2,314.0	2,314.0	2,367.1
Residential	98.3	98.9	90.9	101.5	107.7	106.2	104.5	106.7
Commercial	69.5	50.7	47.5	47.3	54.2	53.1	52.7	53.9
Industrial	313.9	327.4	295.6	297.8	305.5	321.6	318.4	327.3
Transportation	1,410.9	1,562.2	1,598.3	1,655.2	1,702.2	1,685.8	1,715.2	1,731.8
Electricity Generation	101.0	74.3	104.3	96.7	91.0	100.9	77.4	96.3
U.S. Territories	27.4	35.5	35.4	35.2	34.3	46.5	45.7	51.2
Geothermal*	0.4	0.4	0.4	0.4	0.4	0.4	0.4	0.3
Total	4,711.7	5,263.2	5,278.7	5,345.9	5,545.1	5,448.0	5,501.4	5,551.6

NE (Not estimated)

NO (Not occurring)

+ Does not exceed 0.05 Tg CO₂ Eq.

* Although not technically a fossil fuel, geothermal energy-related CO₂ emissions are included for reporting purposes.

Note: Totals may not sum due to independent rounding.

Trends in CO₂ emissions from fossil fuel combustion are influenced by many long-term and short-term factors. On a year-to-year basis, the overall demand for fossil fuels in the United States and other countries generally fluctuates in response to changes in general economic conditions, energy prices, weather, and the availability of non-fossil alternatives. For example, in a year with increased consumption of goods and services, low fuel prices, severe summer and winter weather conditions, nuclear plant closures, and lower precipitation feeding hydroelectric dams, there would likely be proportionally greater fossil fuel consumption than a year with poor economic performance, high fuel prices, mild temperatures, and increased output from nuclear and hydroelectric plants.

Longer-term changes in energy consumption patterns, however, tend to be more a function of aggregate societal trends that affect the scale of consumption (e.g., population, number of cars, and size of houses), the efficiency with which energy is used in equipment (e.g., cars, power plants, steel mills, and light bulbs), and social planning and consumer behavior (e.g., walking, bicycling, or telecommuting to work instead of driving).

Carbon dioxide emissions also depend on the source of energy and its carbon intensity. The amount of carbon in fuels varies significantly by fuel type. For example, coal contains the highest amount of carbon per unit of useful

² An additional discussion of fossil fuel emission trends is presented in the Trends in U.S. Greenhouse Gas Emissions Chapter.

energy. Petroleum has roughly 75 percent of the carbon per unit of energy as coal, and natural gas has only about 55 percent.³ Producing a unit of heat or electricity using natural gas instead of coal can reduce the CO₂ emissions associated with energy consumption, and using nuclear or renewable energy sources (e.g., wind) can essentially eliminate emissions (see Box 3-2). Table 3-4 shows annual changes in emissions during the last six years for coal, petroleum, and natural gas in selected sectors.

Table 3-4: Annual Change in CO₂ Emissions from Fossil Fuel Combustion for Selected Fuels and Sectors (Tg CO₂ Eq. and Percent)

Sector	Fuel Type	1999 to 2000	2000 to 2001	2001 to 2002	2002 to 2003				
Electricity Generation	Coal	87.6	5%	-62.6	-3%	22.2	1%	36.6	2%
Electricity Generation	Natural Gas	20.8	8%	8.4	3%	16.5	6%	-28.0	-9%
Electricity Generation	Petroleum	-5.6	-6%	9.8	11%	-23.5	-23%	18.9	24%
Transportation ^a	Petroleum	47.0	3%	-16.4	-1%	29.4	2%	16.6	1%
Residential	Natural Gas	13.9	5%	-10.7	-4%	6.2	2%	11.5	4%
Commercial	Natural Gas	7.1	4%	-7.9	-5%	4.2	3%	2.0	1%
Industrial	Coal	1.1	1%	-4.4	-3%	-7.9	-6%	0.8	1%
Industrial	Natural Gas	8.2	2%	-38.5	-8%	9.5	2%	-27.6	-6%
All Sectors^b	All Fuels^b	199.2	4%	-97.1	-2%	53.5	1%	50.2	1%

^a Excludes emissions from International Bunker Fuels.

^b Includes fuels and sectors not shown in table.

In the United States, 86 percent of the energy consumed in 2003 was produced through the combustion of fossil fuels such as coal, natural gas, and petroleum (see Figure 3-3 and Figure 3-4). The remaining portion was supplied by nuclear electric power (8 percent) and by a variety of renewable energy sources (6 percent), primarily hydroelectric power and biofuels (EIA 2004a). Specifically, petroleum supplied the largest share of domestic energy demands, accounting for an average of 39 percent of total energy consumption from 1990 through 2003. Natural gas and coal followed in order of importance, accounting for 24 and 23 percent of total consumption, respectively. Petroleum was consumed primarily in the transportation end-use sector, the vast majority of coal was used in electricity generation, and natural gas was broadly consumed in all end-use sectors except transportation (see Figure 3-5) (EIA 2004a).

Figure 3-3: 2003 U.S. Energy Consumption by Energy Source

Figure 3-4: U.S. Energy Consumption (Quadrillion Btu)

Figure 3-5: 2003 CO₂ Emissions from Fossil Fuel Combustion by Sector and Fuel Type

Fossil fuels are generally combusted for the purpose of producing energy for useful heat and work. During the combustion process, the carbon stored in the fuels is oxidized and emitted as CO₂ and smaller amounts of other gases, including CH₄, CO, and NMVOCs.⁴ These other carbon containing non-CO₂ gases are emitted as a by-product of incomplete fuel combustion, but are, for the most part, eventually oxidized to CO₂ in the atmosphere. Therefore, except for the soot and ash left behind during the combustion process, all the carbon in fossil fuels used to produce energy is eventually converted to atmospheric CO₂.

³ Based on national aggregate carbon content of all coal, natural gas, and petroleum fuels combusted in the United States.

⁴ See the sections entitled Stationary Combustion and Mobile Combustion in this chapter for information on non-CO₂ gas emissions from fossil fuel combustion.

[BEGIN BOX]

Box 3-1: Weather and Non-Fossil Energy Effects on CO₂ from Fossil Fuel Combustion Trends

In 2003, weather conditions became cooler in both the winter and summer. Heating degree days in the United States were 2 percent below normal (see Figure 3-6), while cooling degree days in 2003 were 5 percent above normal (see Figure 3-7) (EIA 2004f).⁵ Winter conditions were colder in 2003 than in 2002, which, coupled with a 1.2 percent increase in the U.S. housing stock (EIA 2004f), led to an increase in demand for heating fuels, despite escalating fuel prices for heating fuels. Though the summer of 2003 was cooler than the near record heat of 2002, demand for electricity increased most likely due to the growing economy and increase in housing stock.

Figure 3-6: Annual Deviations from Normal Heating Degree Days for the United States (1949-2003)

Figure 3-7: Annual Deviations from Normal Cooling Degree Days for the United States (1949-2003)

Although no new U.S. nuclear power plants have been constructed in recent years, the utilization (i.e., capacity factors⁶) of existing plants remained high, at 88 percent in 2003. This utilization level actually represents a 2 percent decrease in electricity output by nuclear plants, down from a record high of 90 percent in 2002. Electricity output by hydroelectric power plants increased in 2003 by approximately 4 percent. Nevertheless, electricity generated by nuclear plants in 2003 provided almost 3 times as much of the energy consumed in the United States as hydroelectric plants (EIA 2004a). Aggregate nuclear and hydroelectric power plant capacity factors since 1973 are shown in Figure 3-8.

Figure 3-8: Aggregate Nuclear and Hydroelectric Power Plant Capacity Factors in the United States (1973-2003)

[END BOX]

For the purpose of international reporting, the IPCC (IPCC/UNEP/OECD/IEA 1997) recommends that particular adjustments be made to national fuel consumption statistics. Certain fossil fuels can be manufactured into plastics, asphalt, lubricants, or other products. A portion of the carbon consumed for these non-energy products can be stored (i.e., sequestered) indefinitely. To account for the fact that the carbon in these fuels ends up in products instead of being combusted (i.e., oxidized and released into the atmosphere), consumption of fuels for non-energy purposes is estimated and subtracted from total fuel consumption estimates. Emissions from non-energy uses of fuels are estimated in the Carbon Emitted and Stored in Products from Non-Energy Uses of Fossil Fuels section in this chapter.

⁵ Degree days are relative measurements of outdoor air temperature. Heating degree days are deviations of the mean daily temperature below 65° F, while cooling degree days are deviations of the mean daily temperature above 65° F. Heating degree days have a considerably greater affect on energy demand and related emissions than do cooling degree days. Excludes Alaska and Hawaii. Normals are based on data from 1971 through 2000. The variation in these normals during this time period was ±10 percent and ±14 percent for heating and cooling degree days, respectively (99 percent confidence interval).

⁶ The capacity factor is defined as the ratio of the electrical energy produced by a generating unit for a given period of time to the electrical energy that could have been produced at continuous full-power operation during the same period (EIA 2004a).

According to the UNFCCC reporting guidelines, CO₂ emissions from the consumption of fossil fuels for aviation and marine international transport activities (i.e., international bunker fuels) should be reported separately, and not included in national emission totals. Estimates of international bunker fuel emissions for the United States are provided in Table 3-5.

Table 3-5: CO₂ Emissions from International Bunker Fuels (Tg CO₂ Eq.)*

Vehicle Mode	1990	1997	1998	1999	2000	2001	2002	2003
Aviation	46.2	55.9	56.7	58.9	60.5	59.4	61.8	59.6
Marine	67.3	54.0	57.9	46.4	40.9	38.5	27.7	24.6
Total	113.5	109.9	114.6	105.3	101.4	97.9	89.5	84.2

* See International Bunker Fuels section for additional detail.

Note: Totals may not sum due to independent rounding.

End-Use Sector Consumption

An alternative method of presenting CO₂ emissions is to allocate emissions associated with electricity generation to the sectors in which it is used. Four end-use sectors were defined: industrial, transportation, residential, and commercial.⁷ For the discussion below, electricity generation emissions have been distributed to each end-use sector based upon the sector's share of national electricity consumption. This method of distributing emissions assumes that each sector consumes electricity generated from an equally carbon-intensive mix of fuels and other energy sources. In reality, sources of electricity vary widely in carbon intensity (e.g., coal versus wind power). By giving equal carbon-intensity weight to each sector's electricity consumption, emissions attributed to one end-use sector may be somewhat overestimated, while emissions attributed to another end-use sector may be slightly underestimated. After the end-use sectors are discussed, emissions from electricity generation are addressed separately. Emissions from U.S. territories are also calculated separately due to a lack of end-use-specific consumption data. Table 3-6 and Figure 3-9 summarize CO₂ emissions from direct fossil fuel combustion and prorated electricity generation emissions from electricity consumption by end-use sector.

Table 3-6: CO₂ Emissions from Fossil Fuel Combustion by End-Use Sector (Tg CO₂ Eq.)

End-Use Sector	1990	1997	1998	1999	2000	2001	2002	2003
Transportation	1,449.8	1,606.4	1,636.5	1,693.9	1,741.0	1,723.1	1,755.4	1,770.4
Combustion	1,446.8	1,603.3	1,633.4	1,690.8	1,737.7	1,719.7	1,752.3	1,767.2
Electricity	3.0	3.1	3.1	3.2	3.4	3.4	3.2	3.2
Industrial	1,553.9	1,703.0	1,668.5	1,651.2	1,684.4	1,587.4	1,579.0	1,572.9
Combustion	882.8	963.8	911.6	888.1	905.0	878.2	876.6	858.6
Electricity	671.1	739.2	757.0	763.1	779.4	709.3	702.4	714.3
Residential	924.8	1,040.7	1,044.4	1,063.5	1,124.2	1,116.2	1,145.0	1,168.9
Combustion	339.6	370.6	338.6	359.3	379.1	367.0	371.4	385.1
Electricity	585.3	670.2	705.8	704.2	745.0	749.2	773.6	783.8
Commercial	755.1	876.7	892.9	901.2	959.5	972.7	973.9	983.1
Combustion	224.2	237.2	219.7	222.3	235.2	226.7	230.0	234.0
Electricity	530.9	639.5	673.2	678.9	724.3	745.9	743.9	749.2
U.S. Territories	28.0	36.4	36.3	36.2	35.9	48.6	48.1	56.2
Total	4,711.7	5,263.2	5,278.7	5,345.9	5,545.1	5,448.0	5,501.4	5,551.6
Electricity Generation	1,790.3	2,051.9	2,139.0	2,149.3	2,252.1	2,207.8	2,223.0	2,250.5

Note: Totals may not sum due to independent rounding. Emissions from fossil fuel combustion by electricity generation are allocated based on aggregate national electricity consumption by each end-use sector.

⁷ See Glossary (Annex 6.8) for more detailed definitions of the industrial, residential, commercial, and transportation end-use sector, as well as electricity generation.

Figure 3-9: 2003 End-Use Sector Emissions of CO₂ from Fossil Fuel Combustion

Transportation End-Use Sector

Using this allocation method, the transportation end-use sector accounted for the largest share (approximately 32 percent) of CO₂ emissions from fossil fuel combustion.⁸ Almost all of the energy consumed in the transportation sector was petroleum-based, with nearly two-thirds being gasoline consumption in automobiles and other highway vehicles. Other fuel uses, especially diesel fuel for freight trucks and jet fuel for aircraft, accounted for the remainder.⁹

Carbon dioxide emissions from fossil fuel combustion for transportation increased by 22 percent from 1990 to 2003, to 1,770.4 Tg CO₂ Eq. The growth in transportation end-use sector emissions has been relatively steady, excluding a 4 percent single year increase in 1999 and slight decreases in 1991 and 2001. Like overall energy demand, transportation fuel demand is a function of many short and long-term factors. In the short term only minor adjustments can generally be made through consumer behavior (e.g., not driving as far for summer vacation). However, long-term adjustments such as vehicle purchase choices, transport mode choice and access (i.e., trains versus planes), and urban planning can have a significant impact on fuel demand.

In 2003, CO₂ emissions from the transportation sector increased by approximately 1 percent. A 12 percent increase in the price of motor gasoline in 2003 tempered the effects of the growing economy¹⁰ on demand for vehicle fuel (see Figure 3-10).

Since 1990, travel activity in the United States has grown more rapidly than population, with a 16 percent increase in vehicle miles traveled per capita. In the meantime, improvements in the average fuel efficiency of the U.S. vehicle fleet stagnated after increasing steadily since 1976 (FHWA 1996 through 2004). The average miles per gallon achieved by the U.S. vehicle fleet has remained fairly constant since 1991. This trend is due, in part, to the increasing dominance of new motor vehicle sales by less fuel-efficient light-duty trucks and sport-utility vehicles (see Figure 3-11).

Figure 3-10: Motor Gasoline Retail Prices (Real)

Figure 3-11: Motor Vehicle Fuel Efficiency

Table 3-7 provides a detailed breakdown of CO₂ emissions by fuel category and vehicle type for the transportation end-use sector. Fifty-nine percent of the emissions from this end-use sector in 2003 were the result of the combustion of motor gasoline in passenger cars and light-duty trucks. Diesel highway vehicles and jet aircraft were also significant contributors, accounting for 18 and 12 percent of CO₂ emissions from the transportation end-use sector, respectively.¹¹ For information on transportation-related CO₂ emissions from agriculture and construction equipment, other off-road equipment, and recreational vehicles, please refer to Table 3-36 in Annex 3.2.

Table 3-7: CO₂ Emissions from Fossil Fuel Combustion in Transportation End-Use Sector (Tg CO₂ Eq.)

Fuel/Vehicle Type	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline	955.2	1,042.5	1,072.9	1,099.9	1,105.9	1,111.2	1,138.7	1,143.7

⁸ Note that electricity generation is actually the largest emitter of CO₂ when electricity is not distributed among end-use sectors.

⁹ See Glossary (Annex 6.8) for a more detailed definition of the transportation end-use sector.

¹⁰ Gross domestic product increased 3.1 percent between 2002 and 2003 (BEA 2004).

¹¹ These percentages include emissions from bunker fuels.

Automobiles	605.1	589.8	608.6	618.4	620.1	622.3	636.1	630.2
Light-Duty Trucks	301.0	406.1	416.3	432.6	435.3	438.7	450.9	460.9
Other Trucks	37.7	33.3	34.6	35.7	37.3	37.1	38.7	39.6
Buses	0.3	0.4	0.4	0.4	0.4	0.4	0.3	0.3
Motorcycles	1.7	1.7	1.8	1.8	1.8	1.6	1.6	1.6
Boats (Recreational)	9.4	11.2	11.2	11.1	11.2	11.2	11.1	11.0
Distillate Fuel Oil (Diesel)	265.1	338.4	348.4	362.2	374.0	383.2	378.1	392.6
Automobiles	7.4	5.6	5.2	4.0	3.4	3.5	3.4	3.4
Light-Duty Trucks	10.7	15.1	15.4	16.3	16.6	17.2	16.9	17.6
Other Trucks	178.4	246.2	255.2	268.1	282.7	289.9	288.3	301.1
Buses	7.5	8.6	8.7	9.7	9.4	8.8	8.2	8.0
Locomotives	33.3	37.0	37.4	38.5	38.4	39.4	37.9	39.6
Ships & Boats	16.3	16.9	15.0	17.4	17.3	19.1	18.4	17.0
Ships (Bunkers)	11.4	9.1	11.5	8.2	6.2	5.2	5.1	6.0
Jet Fuel	220.4	232.1	235.6	242.9	251.2	240.4	234.4	228.6
Commercial Aircraft	117.2	128.5	126.3	136.4	140.6	132.8	121.7	122.8
Military Aircraft	34.8	21.0	21.5	20.6	21.0	22.8	20.4	20.5
General Aviation Aircraft	6.3	6.1	7.7	9.2	9.2	9.0	9.3	9.4
Other Aircraft ^b	15.9	20.6	23.4	17.8	19.9	16.4	21.2	16.3
Aircraft (Bunkers)	46.2	55.9	56.7	58.9	60.5	59.4	61.8	59.6
Aviation Gasoline	3.1	2.7	2.4	2.7	2.5	2.4	2.3	2.2
General Aviation Aircraft	3.1	2.7	2.4	2.7	2.5	2.4	2.3	2.2
Residual Fuel Oil	79.3	55.5	52.6	51.9	69.2	45.7	50.4	48.1
Ships & Boats ^c	23.4	10.6	6.2	13.7	34.6	12.4	27.7	29.5
Ships (Bunkers) ^c	55.8	44.9	46.4	38.2	34.6	33.2	22.6	18.6
Natural Gas	35.9	41.1	35.1	35.6	35.5	33.9	37.1	35.4
Automobiles	+	+	+	+	+	+	+	+
Light Trucks	+	+	+	+	+	+	+	+
Buses	+	0.2	0.2	0.3	0.4	0.5	0.6	0.6
Pipeline	35.9	40.9	34.9	35.3	35.0	33.4	36.4	34.8
LPG	1.4	0.8	1.0	0.8	0.7	0.8	0.9	0.8
Light Trucks	0.5	0.4	0.4	0.3	0.3	0.3	0.3	0.3
Other Trucks	0.8	0.4	0.6	0.5	0.4	0.5	0.5	0.5
Buses	+	+	+	+	+	+	+	+
Electricity	3.0	3.1	3.1	3.2	3.4	3.4	3.2	3.2
Rail	3.0	3.1	3.1	3.2	3.4	3.4	3.2	3.2
Total (Including Bunkers)^d	1,563.3	1,716.2	1,751.1	1,799.2	1,842.5	1,821.0	1,844.9	1,854.6
Total (Excluding Bunkers)^d	1,449.8	1,606.4	1,636.5	1,693.9	1,741.0	1,723.1	1,755.4	1,770.4

Note: Totals may not sum due to independent rounding.

^a Emissions are no longer allocated to gasoline and diesel consumption from agriculture and construction, and electricity consumption from pipelines. This based on recognition that EIA statistics account for these activities in the industrial sector.

^b This category represents all other jet fuel consumption, and may include some small commercial aircraft and jet fuel used for heating oil.

^c Fluctuations in emission estimates from the combustion of residual fuel oil are currently unexplained, but may be related to data collection problems.

^d Official estimates exclude emissions from the combustion of both aviation and marine international bunker fuels; however, estimates including international bunker fuel-related emissions are presented for informational purposes.

+ Less than 0.05 Tg CO₂ Eq.

Industrial End-Use Sector

The industrial end-use sector accounted for 28 percent of CO₂ emissions from fossil fuel combustion. On average, 55 percent of these emissions resulted from the direct consumption of fossil fuels for steam and process heat production. The remaining 45 percent was associated with their consumption of electricity for uses such as motors, electric furnaces, ovens, and lighting.

The industrial end-use sector includes activities such as manufacturing, construction, mining, and agriculture.¹² The largest of these activities in terms of energy consumption is manufacturing, which was estimated in 1998 to have accounted for about 84 percent of industrial energy consumption (EIA 2001a). Just six industries—Petroleum, Chemicals, Primary Metals, Paper, Food, and Stone, Clay, and Glass products—represent 83 percent of total manufacturing energy use.

In theory, emissions from the industrial end-use sector should be highly correlated with economic growth and industrial output, but heating of industrial buildings and agricultural energy consumption is also affected by weather conditions.¹³ In addition, structural changes within the U.S. economy that lead to shifts in industrial output away from energy intensive manufacturing products to less energy intensive products (e.g., from steel to computer equipment) also have a significant affect on industrial emissions.

From 2002 to 2003, total industrial production and manufacturing output increased slightly, by 0.2 and 0.1 percent, respectively (FRB 2004). Also from 2002 to 2003, output increased for the Petroleum Refinery, Nonmetallic Mineral Product, and Chemical industries, but declined for the Primary Metal, Food, and Paper industries (see Figure 3-12).

Figure 3-12: Industrial Production Indexes (Index 1997=100)

Despite the growth in industrial output (44 percent) and the overall U.S. economy (46 percent) from 1990 to 2003, emissions from the industrial end-use sector increased only slightly (by 1 percent). The reasons for the disparity between rapid growth in industrial output and stagnant growth in industrial emissions are not entirely clear. It is likely, though, that several factors have influenced industrial emission trends, including: 1) more rapid growth in output from less energy-intensive industries relative to traditional manufacturing industries, 2) improvements in energy efficiency; and 3) a lowering of the carbon intensity of fossil fuel consumption as industry shifts from its historical reliance on coal and coke to heavier usage of natural gas. In 2003, carbon dioxide emissions from fossil fuel combustion and electricity use within the industrial end-use sectors were 1,572.9 Tg CO₂ Eq., or 0.4 percent below 2002 emissions.

Residential and Commercial End-Use Sectors

The residential and commercial end-use sectors accounted for an average 21 and 18 percent, respectively, of CO₂ emissions from fossil fuel combustion. Both end-use sectors were heavily reliant on electricity for meeting energy needs, with electricity consumption for lighting, heating, air conditioning, and operating appliances contributing to about 67 and 76 percent of emissions from the residential and commercial end-use sectors, respectively. The remaining emissions were largely due to the direct consumption of natural gas and petroleum products, primarily for heating and cooking needs. Coal consumption was a minor component of energy use in both of these end-use sectors. In 2003, CO₂ emissions from fossil fuel combustion and electricity use within the residential and commercial end-use sectors were 1,168.9 Tg CO₂ Eq. and 983.1 Tg CO₂ Eq., respectively.

Emissions from the residential and commercial sectors have generally been increasing since 1990, and are often correlated with short-term fluctuations in energy consumption caused by weather conditions, rather than prevailing economic conditions (see Table 3-6). In the long-term, both end-use sectors are also affected by population growth, regional migration trends, and changes in housing and building attributes (e.g., size and insulation).

¹² See Glossary (Annex 6.8) for a more detailed definition of the industrial end-use sector.

¹³ Some commercial customers are large enough to obtain an industrial price for natural gas and/or electricity and are consequently grouped with the industrial end-use sector in U.S. energy statistics. These misclassifications of large commercial customers likely cause the industrial end-use sector to appear to be more sensitive to weather conditions.

Emissions from natural gas consumption represent over 70 percent of the direct (not including electricity) fossil fuel emissions from the residential and commercial sectors. In 2003, these emissions increased by 4 and 1 percent, respectively, in each of these sectors. Colder winter conditions in the United States (see Figure 3-13) and an increasing U.S. housing stock led to higher demand for natural gas, despite drastic increases in natural gas prices (66 percent).

Figure 3-13: Heating Degree Days¹⁴

Electricity sales to the residential and commercial end-use sectors in 2003 increased by 1 and 0.4 percent, respectively. This trend can largely be attributed to the growing economy (3.1 percent) and increase in U.S. housing stock (1.2 percent), which led to increased demand for electricity. Increased consumption due to these factors was somewhat offset by decreases in air conditioning-related electricity consumption expected with the cooler summer (see Figure 3-14), and increases in electricity prices (1 and 2 percent higher to the residential and commercial sectors, respectively). Electricity-related emissions in both sectors rose with increasing consumption and the higher carbon intensity of electricity generation. Total emissions from the residential sector increased by 2.1 percent in 2003, with emission from the commercial sector 0.9 percent higher than in 2002.

Figure 3-14: Cooling Degree Days¹⁵

Electricity Generation

The process of generating electricity is the single largest source of CO₂ emissions in the United States (39 percent). Electricity was consumed primarily in the residential, commercial, and industrial end-use sectors for lighting, heating, electric motors, appliances, electronics, and air conditioning (see Figure 3-15). Electricity generation also accounted for the largest share of CO₂ emissions from fossil fuel combustion, approximately 41 percent in 2003.

Figure 3-15: Electricity Generation Retail Sales by End-Use Sector

The electric power industry includes all power producers, consisting of both regulated utilities and nonutilities (e.g. independent power producers, qualifying cogenerators, and other small power producers). While utilities primarily generate power for the U.S. electric grid for sale to retail customers, nonutilities produce electricity for their own use, to sell to large consumers, or to sell on the wholesale electricity market (e.g., to utilities for distribution and resale to customers). However, the electric power industry in the United States has undergone significant changes as both federal and state government agencies have modified regulations to create a more competitive market for electricity generation. These changes have led to the growth of nonutility power producers, including the sale of generating capacity by electric utilities to nonutilities. As a result, the Department of Energy no longer categorizes electric power generation into these ownership groups, and instead uses three functional categories: the electric power sector, the commercial sector, and the industrial sector. The electric power sector consists of electric utilities and independent power producers whose primary business is the production of electricity, while the other sectors consist of those producers that indicate their primary business is other than the production of electricity.

In 2003, the amount of electricity generated decreased very slightly, by 0.3 percent. Although total U.S. electricity use actually increased by 1 percent, net generation declined due to increased net imports and reduced losses of

¹⁴ Degree days are relative measurements of outdoor air temperature. Heating degree days are deviations of the mean daily temperature below 65° F. Excludes Alaska and Hawaii. Normals are based on data from 1971 through 2000.

¹⁵ Degree days are relative measurements of outdoor air temperature. Cooling degree days are deviations of the mean daily temperature above 65° F. Excludes Alaska and Hawaii. Normals are based on data from 1971 through 2000.

electricity.¹⁶ However, CO₂ emissions increased by 1.2 percent, as escalating natural gas prices caused power producers to switch from natural gas to coal, a more carbon-intensive fuel. Coal consumption for electricity generation increased by 2.0 percent in 2003, while natural gas consumption decreased by 9.2 percent. As a result of this shift, carbon intensity from energy consumption for electricity generation increased in 2003 (see Table 3-9). Coal is consumed primarily by the electric power sector in the United States, which accounted for 93 percent of total coal consumption for energy purposes in 2003. Electricity generation by nuclear and renewable resources remained relatively stable, increasing 1 percent in 2003.

[BEGIN BOX]

Box 3-2: Carbon Intensity of U.S. Energy Consumption

Fossil fuels are the dominant source of energy in the United States, and CO₂ is emitted as a product from their combustion. Useful energy, however, can be generated from many other sources that do not emit CO₂ in the energy conversion process. In the United States, useful energy is also produced from renewable (i.e., hydropower, biofuels, geothermal, solar, and wind) and nuclear sources.¹⁷

Energy-related CO₂ emissions can be reduced by not only lowering total energy consumption (e.g., through conservation measures) but also by lowering the carbon intensity of the energy sources employed (e.g., fuel switching from coal to natural gas). The amount of carbon emitted from the combustion of fossil fuels is dependent upon the carbon content of the fuel and the fraction of that carbon that is oxidized.¹⁸ Fossil fuels vary in their average carbon content, ranging from about 53 Tg CO₂ Eq./QBtu for natural gas to upwards of 95 Tg CO₂ Eq./QBtu for coal and petroleum coke.¹⁹ In general, the carbon content per unit of energy of fossil fuels is the highest for coal products, followed by petroleum, and then natural gas. Other sources of energy, however, may be directly or indirectly carbon neutral (i.e., 0 Tg CO₂ Eq./Btu). Energy generated from nuclear and many renewable sources do not result in direct emissions of CO₂. Biofuels such as wood and ethanol are also considered to be carbon neutral; although these fuels do emit CO₂, in the long run the CO₂ emitted from biomass consumption does not increase atmospheric CO₂ concentrations if the biogenic carbon emitted is offset by the growth of new biomass.²⁰ The overall carbon intensity of the U.S. economy is thus dependent upon the quantity and combination of fuels and other energy sources employed to meet demand.

Table 3-8 provides a time series of the carbon intensity for each sector of the U.S. economy. The time series incorporates only the energy consumed from the direct combustion of fossil fuels in each sector. For example, the carbon intensity for the residential sector does not include the energy from or emissions related to the consumption of electricity for lighting or wood for heat. Looking only at this direct consumption of fossil fuels, the residential sector exhibited the lowest carbon intensity, which is related to the large percentage of its energy derived from

¹⁶ EIA statistics track net generation, imports, exports, and compare these with the end use of electricity. The difference between these values is classified as “T&D losses and Unaccounted for”. T&D losses refer to electricity lost during the transmission and distribution of electricity from the source to the end user. There also exists a small amount of electricity not accounted for due to data collection time frame differences and nonsampling error.

¹⁷ Small quantities of CO₂, however, are released from some geologic formations tapped for geothermal energy. These emissions are included with fossil fuel combustion emissions from the electricity generation. Carbon dioxide emissions may also be generated from upstream activities (e.g., manufacture of the equipment) associated with fossil fuel and renewable energy activities, but are not accounted for here.

¹⁸ Generally, more than 97 percent of the carbon in fossil fuel is oxidized to CO₂ with most carbon combustion technologies used in the United States.

¹⁹ One exajoule (EJ) is equal to 10¹⁸ joules or 0.9478 QBtu.

²⁰ Net carbon fluxes from changes in biogenic carbon reservoirs in wooded or croplands are accounted for in the estimates for Land-Use Change and Forestry.

natural gas for heating. The carbon intensity of the commercial sector has predominantly declined since 1990 as commercial businesses shift away from petroleum to natural gas. The industrial sector was more dependent on petroleum and coal than either the residential or commercial sectors, and thus had higher carbon intensities over this period. The carbon intensity of the transportation sector was closely related to the carbon content of petroleum products (e.g., motor gasoline and jet fuel, both around 70 Tg CO₂ Eq./EJ), which were the primary sources of energy. Lastly, the electricity generation sector had the highest carbon intensity due to its heavy reliance on coal for generating electricity.

Table 3-8: Carbon Intensity from Direct Fossil Fuel Combustion by Sector (Tg CO₂ Eq./QBtu)

Sector	1990	1997	1998	1999	2000	2001	2002	2003
Residential ^a	57.0	56.5	56.5	56.6	56.6	56.7	56.6	56.5
Commercial ^a	59.3	57.4	57.1	57.1	57.2	57.3	57.2	57.3
Industrial ^a	66.1	65.8	65.2	65.1	65.1	64.7	64.5	65.0
Transportation ^a	70.3	70.1	70.1	70.2	70.3	70.3	70.3	70.3
Electricity Generation ^b	85.8	85.9	85.2	84.9	84.7	84.3	84.1	84.8
U.S. Territories ^c	73.3	73.3	73.2	73.0	72.5	72.9	73.0	73.5
All Sectors^c	72.6	72.5	72.6	72.5	72.5	72.3	72.2	72.5

^a Does not include electricity or renewable energy consumption.

^b Does not include electricity produced using nuclear or renewable energy.

^c Does not include nuclear or renewable energy consumption.

Note: Excludes non-energy fuel use emissions and consumption.

In contrast to Table 3-8, Table 3-9 presents carbon intensity values that incorporate energy consumed from all sources (i.e., fossil fuels, renewables, and nuclear). In addition, the emissions related to the generation of electricity have been attributed to both electricity generation and the end-use sectors in which that electricity was eventually consumed.²¹ This table, therefore, provides a more complete picture of the actual carbon intensity of each end-use sector per unit of energy consumed. The transportation end-use sector in Table 3-9 emerges as the most carbon intensive when all sources of energy are included, due to its almost complete reliance on petroleum products and relatively minor amount of biomass-based fuels such as ethanol. The “other end-use sectors” (i.e., residential, commercial, and industrial) use significant quantities of biofuels such as wood, thereby lowering the overall carbon intensity. The carbon intensity of the electricity generation sector differs greatly from the scenario in Table 3-8, where only the energy consumed from the direct combustion of fossil fuels was included. This difference is due almost entirely to the inclusion of electricity generation from nuclear and hydropower sources, which do not emit CO₂.

Table 3-9: Carbon Intensity from all Energy Consumption by Sector (Tg CO₂ Eq./QBtu)

Sector	1990	1997	1998	1999	2000	2001	2002	2003
Transportation ^a	70.1	69.8	69.8	69.8	69.9	69.8	69.8	69.6
Other End-Use Sectors ^{a, b}	57.8	57.8	57.9	57.3	58.0	58.2	57.5	58.0
Electricity Generation ^c	58.4	58.9	59.1	58.2	59.2	59.3	58.3	59.0
All Sectors^d	61.1	61.0	61.1	60.8	61.3	61.5	60.9	61.2

^a Includes electricity (from fossil fuel, nuclear, and renewable sources) and direct renewable energy consumption.

^b Other End-Use Sectors includes the residential, commercial, and industrial sectors.

^c Includes electricity generation from nuclear and renewable sources.

^d Includes nuclear and renewable energy consumption.

Note: Excludes non-energy fuel use emissions and consumption.

²¹ In other words, the emissions from the generation of electricity are intentionally double counted by attributing them both to electricity generation and the end-use sector in which electricity consumption occurred.

By comparing the values in Table 3-8 and Table 3-9, a few observations can be made. The use of renewable and nuclear energy sources has resulted in a significantly lower carbon intensity of the U.S. economy. Over the thirteen-year period of 1990 through 2003, however, the carbon intensity of U.S. energy consumption has been fairly constant, as the proportion of renewable and nuclear energy technologies has not changed significantly.

The carbon intensity of total energy consumption in the United States has remained fairly constant. Per capita energy consumption has fluctuated, but has generally demonstrated a constant overall trend since 1990 (see Figure 3-16). Due to a general shift from a manufacturing-based economy to a service-based economy, as well as overall increases in efficiency, energy consumption and energy-related CO₂ emissions per dollar of gross domestic product (GDP) have both declined since 1990.

Figure 3-16: U.S. Energy Consumption and Energy-Related CO₂ Emissions Per Capita and Per Dollar GDP

Carbon intensity estimates were developed using nuclear and renewable energy data from EIA (2004a) and fossil fuel consumption data as discussed above and presented in Annex 2.1.

[END BOX]

Methodology

The methodology used by the United States for estimating CO₂ emissions from fossil fuel combustion is conceptually similar to the approach recommended by the IPCC for countries that intend to develop detailed, sectoral-based emission estimates (IPCC/UNEP/OECD/IEA 1997). A detailed description of the U.S. methodology is presented in Annex 2.1, and is characterized by the following steps:

1. *Determine total fuel consumption by fuel type and sector.* Total fossil fuel consumption for each year is estimated by aggregating consumption data by end-use sector (e.g., commercial, industrial, etc.), primary fuel type (e.g., coal, petroleum, gas), and secondary fuel category (e.g., motor gasoline, distillate fuel oil, etc.). Fuel consumption data for the United States were obtained directly from the Energy Information Administration (EIA) of the U.S. Department of Energy (DOE), primarily from the *Monthly Energy Review* and unpublished supplemental tables on petroleum product detail (EIA 2004b). The United States does not include territories in its national energy statistics, so fuel consumption data for territories were collected separately from Grillot (2004).²²

For consistency of reporting, the IPCC has recommended that countries report energy data using the International Energy Agency (IEA) reporting convention and/or IEA data. Data in the IEA format are presented "top down"—that is, energy consumption for fuel types and categories are estimated from energy production data (accounting for imports, exports, stock changes, and losses). The resulting quantities are referred to as "apparent consumption." The data collected in the United States by EIA, and used in this inventory, are, instead, "bottom up" in nature. In other words, they are collected through surveys at the point of delivery or use and aggregated to determine national totals.²³

It is also important to note that U.S. fossil fuel energy statistics are generally presented using gross calorific values (GCV) (i.e., higher heating values). Fuel consumption activity data presented here have not been

²² Fuel consumption by U.S. territories (i.e. American Samoa, Guam, Puerto Rico, U.S. Virgin Islands, Wake Island, and other U.S. Pacific Islands) is included in this report and contributed emissions of 56 Tg CO₂ Eq. in 2003.

²³ See IPCC Reference Approach for estimating CO₂ emissions from fossil fuel combustion in Annex 4 for a comparison of U.S. estimates using top-down and bottom-up approaches.

adjusted to correspond to international standard, which are to report energy statistics in terms of net calorific values (NCV) (i.e., lower heating values).²⁴

2. *Subtract uses accounted for in the Industrial Processes chapter.* Portions of the fuel consumption data for five fuel categories—coking coal, petroleum coke, natural gas, residual fuel oil, and other oil—were reallocated to the industrial processes chapter, as they were consumed during non-energy related industrial activity. To make these adjustments, additional data were collected from Gambogi (2004), EFMA (1995), U.S. Census Bureau (1991 through 1994), U.S. Census Bureau (1998 through 2003), U.S. Census Bureau (2003), U.S. Census Bureau (2004a), EIA (2000 through 2004), EIA (2001b), USGS (2003 through 2004), USGS (1998 through 2002), USGS (1995), USGS (1995 through 2004), USGS (1991 through 1994), USGS (1991 through 2003), U.S. International Trade Commission (2004a), U.S. International Trade Commission (2004b), and Onder and Bagdoyan (1993).²⁵
3. *Adjust for biofuels, conversion of fossil fuels, and exports of CO₂.* Fossil fuel consumption estimates are adjusted downward to exclude 1) fuels with biogenic origins, 2) fuels created from other fossil fuels, and 3) exports of CO₂. Fuels with biogenic origins are assumed to result in no net CO₂ emissions, and must be subtracted from fuel consumption estimates. These fuels include ethanol added to motor gasoline and biomass gas used as natural gas. Synthetic natural gas is created from industrial coal, and is currently included in EIA statistics for both coal and natural gas. Therefore, synthetic natural gas is subtracted from energy consumption statistics.²⁶ Since October 2000, the Dakota Gasification Plant has been exporting CO₂ to Canada by pipeline. Since this CO₂ is not emitted to the atmosphere in the United States, energy used to produce this CO₂ is subtracted from energy consumption statistics. To make these adjustments, additional data for ethanol and biogas were collected from EIA (2004b) and data for synthetic natural gas were collected from EIA (2004e), and data for CO₂ exports were collected from the Dakota Gasification Company (2003), Fitzpatrick (2002), and Erickson (2003).
4. *Adjust for fuels consumed for non-energy uses.* U.S. aggregate energy statistics include consumption of fossil fuels for non-energy purposes. Depending on the end-use, this can result in storage of some or all of the carbon contained in the fuel for a period of time. As the emission pathways of carbon used for non-energy purposes are vastly different than fuel combustion, these emissions are estimated separately in the Carbon Emitted and Stored in Products from Non-Energy Uses of Fossil Fuels section in this chapter. Therefore, the amount of fuels used for non-energy purposes was subtracted from total fuel consumption. Data on non-fuel consumption was provided by EIA (2004b).
5. *Subtract consumption of international bunker fuels.* According to the UNFCCC reporting guidelines emissions from international transport activities, or bunker fuels, should not be included in national totals. U.S. energy consumption statistics include these bunker fuels (e.g., distillate fuel oil, residual fuel oil, and jet fuel) as part of consumption by the transportation end-use sector, however, so emissions from international transport activities were calculated separately following the same procedures used for emissions from consumption of all fossil fuels (i.e., estimation of consumption, determination of carbon content, and adjustment for the fraction of carbon not oxidized).²⁷ The Office of the Under Secretary of Defense (Installations and Environment) and the Defense Energy Support Center (Defense Logistics Agency) of the U.S. Department of Defense (DoD) (DESC 2004) supplied data on military jet fuel use. Commercial jet fuel use was obtained from BEA (1991 through

²⁴ A crude convention to convert between gross and net calorific values is to multiply the heat content of solid and liquid fossil fuels by 0.95 and gaseous fuels by 0.9 to account for the water content of the fuels. Biomass-based fuels in U.S. energy statistics, however, are generally presented using net calorific values.

²⁵ See sections on Iron and Steel Production, Ammonia Manufacture, Petrochemical Production, Titanium Dioxide Production, Ferroalloy Production, and Aluminum Production in the Industrial Processes chapter.

²⁶ These adjustments are explained in greater detail in Annex 2.1.

²⁷ See International Bunker Fuels section in this chapter for a more detailed discussion.

2004) and DOT (1991 through 2004); residual and distillate fuel use for civilian marine bunkers was obtained from DOC (1991 through 2004). Consumption of these fuels was subtracted from the corresponding fuels in the transportation end-use sector. Estimates of international bunker fuel emissions are discussed further in the section entitled International Bunker Fuels.

6. *Determine the total carbon content of fuels consumed.* Total carbon was estimated by multiplying the amount of fuel consumed by the amount of carbon in each fuel. This total carbon estimate defines the maximum amount of carbon that could potentially be released to the atmosphere if all of the carbon in each fuel was converted to CO₂. The carbon content coefficients used by the United States were obtained from EIA's *Emissions of Greenhouse Gases in the United States 2003* (EIA 2004c) and EIA's *Monthly Energy Review* and unpublished supplemental tables on petroleum product detail EIA (EIA 2004b). They are presented in Annexes 2.1 and 2.2.
7. *Adjust for carbon that does not oxidize during combustion.* Because combustion processes are not 100 percent efficient, some of the carbon contained in fuels is not emitted to the atmosphere. Rather, it remains behind as soot and ash. The estimated amount of carbon not oxidized due to inefficiencies during the combustion process was assumed to be 1 percent for petroleum and coal and 0.5 percent for natural gas (see Annex 2.1). Unoxidized or partially oxidized organic (i.e., carbon containing) combustion products were assumed to have eventually oxidized to CO₂ in the atmosphere.²⁸ IPCC provided fraction oxidized values for petroleum and natural gas (IPCC/UNEP/OECD/IEA 1997). Bechtel (1993) provided the fraction oxidation value for coal.
8. *Allocate transportation emissions by vehicle type.* This report provides a more detailed accounting of emissions from transportation because it is such a large consumer of fossil fuels in the United States.²⁹ For fuel types other than jet fuel, fuel consumption data by vehicle type and transportation mode were used to allocate emissions by fuel type calculated for the transportation end-use sector. For jet fuel, CO₂ emissions were calculated directly based on reported consumption of fuel. For highway vehicles, annual estimates of combined motor gasoline and diesel fuel consumption by vehicle category were obtained from FHWA (1996 through 2004); for each vehicle category, the percent gasoline, diesel, and other (e.g., CNG, LPG) fuel consumption are estimated using data from DOE (1993 through 2004). For non-highway vehicles, activity data were obtained from AAR (2004), BEA (1991 through 2004), Benson (2002 through 2004), DOE (1993 through 2004), DESC (2004), DOC (1991 through 2004), DOT (1991 through 2004), EIA (2002a), EIA (2002b), EIA (2004a), EIA (2004b), EIA (2003 through 2004), EIA (1991 through 2004), EPA (2004c), and FAA (2004). Heat contents and densities were obtained from EIA (2004a) and USAF (1998).³⁰ The difference between total U.S. jet fuel consumption (as reported by EIA) and civilian air carrier consumption for both domestic and international flights (as reported by DOT and BEA) plus military jet fuel consumption is reported as "other" under the jet fuel category in Table 3-7, and includes such fuel uses as blending with heating oils and fuel used for chartered aircraft flights.

Uncertainty

For estimates of CO₂ from fossil fuel combustion, the amount of CO₂ emitted is directly related to the amount of fuel consumed, the fraction of the fuel that is oxidized, and the carbon content of the fuel. Therefore, a careful accounting of fossil fuel consumption by fuel type, average carbon contents of fossil fuels consumed, and production of fossil fuel-based products with long-term carbon storage should yield an accurate estimate of CO₂ emissions.

²⁸ See Indirect CO₂ from CH₄ Oxidation section in this chapter for a more detailed discussion.

²⁹ Electricity generation is not considered a final end-use sector, because energy is consumed primarily to provide electricity to the other sectors.

³⁰ For a more detailed description of the data sources used for the analysis of the transportation end use sector see the Mobile Combustion (excluding CO₂) and International Bunker Fuels sections of the Energy chapter, Annex 3.2, and Annex 3.7.

Nevertheless, there are uncertainties in the consumption data, carbon content of fuels and products, and carbon oxidation efficiencies. For example, given the same primary fuel type (e.g., coal, petroleum, or natural gas), the amount of carbon contained in the fuel per unit of useful energy can vary. For the United States, however, the impact of these uncertainties on overall CO₂ emission estimates is believed to be relatively small. See, for example, Marland and Pippin (1990).

Although statistics of total fossil fuel and other energy consumption are relatively accurate, the allocation of this consumption to individual end-use sectors (i.e., residential, commercial, industrial, and transportation) is less certain. For example, for some fuels the sectoral allocations are based on price rates (i.e., tariffs), but a commercial establishment may be able to negotiate an industrial rate or a small industrial establishment may end up paying an industrial rate, leading to a misallocation of emissions. Also, the deregulation of the natural gas industry and the more recent deregulation of the electric power industry have likely led to some minor problems in collecting accurate energy statistics as firms in these industries have undergone significant restructuring.

To calculate the total CO₂ emission estimate from energy-related fossil fuel combustion, the amount of fuels used in these non-energy production processes were subtracted from the total fossil fuel consumption for 2003. The amount of CO₂ emissions resulting from non-energy related fossil fuel use has been calculated separately and reported in the Carbon Emitted from Non-Energy Uses of Fossil Fuels section of this report. Additionally, inefficiencies in the combustion process, which can result in ash or soot remaining unoxidized for long periods, were also assumed. These factors all contribute to the uncertainty in the CO₂ estimates. Detailed discussions on the uncertainties associated with Carbon emitted from Non-Energy Uses of Fossil Fuels can be found within that section of this chapter.

Various sources of uncertainty surround the estimation of emissions from international bunker fuels, which are subtracted from the U.S. totals (see the detailed discussions on these uncertainties provided in the International Bunker Fuels section of this chapter). Another source of uncertainty is fuel consumption by U.S. territories. The United States does not collect energy statistics for its territories at the same level of detail as for the fifty states and the District of Columbia. Therefore, estimating both emissions and bunker fuel consumption by these territories is difficult.

Uncertainties in the emission estimates presented above also result from the data used to allocate CO₂ emissions from the transportation end-use sector to individual vehicle types and transport modes. In many cases, bottom-up estimates of fuel consumption by vehicle type do not match aggregate fuel-type estimates from EIA. Further research is planned to improve the allocation into detailed transportation end-use sector emissions. In particular, residual fuel consumption data for marine vessels are highly uncertain, as shown by the large fluctuations in emissions that do not mimic changes in other variables such as shipping ton miles.

The uncertainty analysis was performed by primary fuel type for each end-use sector, using the IPCC-recommended Tier 2 uncertainty estimation methodology, Monte Carlo Simulation technique, with @RISK software. For this uncertainty estimation, the inventory estimation model for CO₂ from fossil fuel combustion was integrated with the relevant inventory variables from the inventory estimation model for International Bunker Fuels, to realistically characterize the interaction (or endogenous correlation) between the variables of these two models. About 150 input variables were modeled for CO₂ from energy-related Fossil Fuel Combustion (including about 10 for non-energy fuel consumption and about 20 for International Bunker Fuels).

In developing the uncertainty estimation model, uniform distributions were assumed for all activity-related input variables and emission factors, based on the SAIC/EIA (2001) report.³¹ Triangular distributions were assigned for

³¹ SAIC/EIA (2001) characterizes the underlying probability density function for the input variables as a combination of uniform and normal distributions (the former to represent the bias component and the latter to represent the random component). However, for purposes of the current uncertainty analysis, it was determined that uniform distribution was more appropriate to characterize the probability density function underlying each of these variables.

the oxidization factors (or combustion efficiencies). The uncertainty ranges were assigned to the input variables based on the data reported in SAIC/EIA (2001) and on conversations with various agency-personnel.³²

The uncertainty ranges for the activity-related input variables were typically asymmetric around their inventory estimates; the uncertainty ranges for the emissions factors were symmetric. Bias (or systematic uncertainties) associated with these variables accounted for much of the uncertainties associated with these variables (SAIC/EIA 2001).³³ For purposes of this uncertainty analysis, each input variable was simulated 10,000 times through Monte Carlo Sampling.

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-10. Fossil fuel combustion CO₂ emissions in 2003 were estimated to be between 5,474.3 and 5,863.3 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 1 percent below to 6 percent above the 2003 emission estimate of 5,551.6 Tg CO₂ Eq.

Table 3-10: Tier 2 Quantitative Uncertainty Estimates for CO₂ Emissions from Energy-related Fossil Fuel Combustion by Fuel Type and Sector (Tg CO₂ Eq. and Percent)

Fuel/Sector	2003 Emission		Uncertainty Range Relative to Emission Estimate ^a			
	Estimate (Tg CO ₂ Eq.)		Relative to Emission Estimate ^a		Relative to Emission Estimate ^a	
			(Tg CO ₂ Eq.)	(%)	(Tg CO ₂ Eq.)	(%)
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Coal^b	2,013.8	1,962.0	2,220.5	-3%	+10%	
Residential	1.1	1.1	1.3	-5%	+16%	
Commercial	9.3	8.9	10.8	-4%	+16%	
Industrial	123.4	120.0	144.1	-3%	+17%	
Transportation	NE	NE	NE	NA	NA	
Electricity Generation	1,876.3	1815.7	2,072.5	-3%	+10%	
U.S. Territories	3.6	3.2	4.3	-12%	+20%	
Natural Gas^b	1,170.3	1,161.0	1,229.9	-1%	+5%	
Residential	277.3	270.5	297.7	-2%	+7%	
Commercial	170.7	166.5	183.2	-2%	+7%	
Industrial	407.9	396.6	439.0	-3%	+8%	
Transportation	35.4	34.5	38.0	-2%	+7%	
Electricity Generation	277.6	270.5	292.8	-3%	+5%	
U.S. Territories	1.4	1.2	1.7	-12%	+17%	
Petroleum^b	2,367.1	2,256.9	2,518.5	-5%	+6%	
Residential	106.7	101.7	112.5	-5%	+6%	
Commercial	53.9	51.6	56.7	-4%	+5%	
Industrial	327.3	283.4	386.0	-13%	+18%	
Transportation	1,731.8	1,627.9	1,857.2	-6%	+7%	
Electric Utilities	96.3	93.8	101.8	-3%	+6%	
U.S. Territories	51.2	47.6	57.2	-7%	+12%	
Total (excluding Geothermal)^b	5,551.2	5,474.0	5,863.0	-1%	+6%	

³² In the SAIC/EIA (2001) report, the quantitative uncertainty estimates were developed for each of the three major fossil fuels used within each end-use sector; the variations within the sub-fuel types within each end-use sector were not modeled. However, for purposes of assigning uncertainty estimates to the sub-fuel type categories within each end-use sector in the current uncertainty analysis, SAIC/EIA (2001)-reported uncertainty estimates were extrapolated.

³³ Although, in general, random uncertainties are the main focus of statistical uncertainty analysis, when the uncertainty estimates are elicited from experts, their estimates include both random and systematic uncertainties. Hence, both these types of uncertainties are represented in this uncertainty analysis.

Geothermal	0.3	NE	NE	NE	NE
Total (including Geothermal)^{b,c}	5,551.6	5,474.3	5,863.3	-1%	+6%

NA (Not Applicable)

NE (Not Estimated)

Notes:

^aRange of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

^bThe low and high estimates for total emissions were calculated separately through simulations and, hence, the low and high emission estimates for the sub-source categories do not sum to total emissions.

^cGeothermal emissions added for reporting purposes, but an uncertainty analysis was not performed for CO₂ emissions from geothermal production.

QA/QC and Verification

A source-specific QA/QC plan for CO₂ from fossil fuel combustion was developed and implemented. This effort included a Tier 1 analysis, as well as portions of a Tier 2 analysis. The Tier 2 procedures that were implemented involved checks specifically focusing on the activity data and methodology used for estimating CO₂ emissions from fossil fuel combustion in the United States. Emission totals for the different sectors and fuels were compared and trends were investigated to determine whether any corrective actions were needed. Minor corrective actions were taken.

Recalculations Discussion

A major change this year was the decision to report emissions from fuels used for non-energy purposes separately from fuel combustion emission estimates. Previously, the carbon stored from non-energy uses was subtracted from total carbon in fuels. However, this method resulted in non-energy emissions being reported within the emission estimates for fuel combustion. This year, these emissions are reported separately in the section entitled "Carbon Emitted and Stored in Products from Non-Energy Uses of Fossil Fuels."

Bunker fuel consumption estimates are now subtracted out from total fuel consumption, instead of subtracting the total carbon in bunker fuels from total potential carbon. This change in methodology does not have any effect on emission estimates, though it allows for clearer and more transparent emission calculations.

Adjustments are now made to the consumption data for residual oil and other oils (>401 deg F) to subtract consumption for carbon black production, for which emissions are estimated in the Petrochemical Production section of the Industrial Processes chapter.

EIA statistics report consumption for non-energy use for a number of fuels to be roughly equivalent to total use of these fuels, though for certain years these data series did not match exactly. After discussions with EIA, it was assumed that 100 percent of the use of these fuels should be assumed to be for non-energy purposes. These fuels are asphalt & road oil, lubricants, naphtha, other oil (>401 deg. F), special naphtha, waxes, and miscellaneous products.

The Energy Information Administration (EIA 2004a) updated energy consumption data for all years. These revisions primarily impacted the emission estimates for 2002.

The combination of the methodological and historical data changes, as well as changes in the estimates of Carbon Emitted and Stored in Products from Non-Energy Uses of Fossil Fuels and International Bunker Fuels (which affect emissions from this source) resulted in an average annual decrease of 115.8 Tg CO₂ Eq. (2.2 percent) in CO₂ emissions from fossil fuel combustion for the period 1990 through 2002. This decrease is largely a result of reporting emissions from non-energy uses separately this year.

Planned Improvements

Several items are being evaluated to improve the estimates of CO₂ emissions from fossil fuel combustion and to reduce uncertainty:

- Currently, the IPCC guidelines recommend a default factor of 0.99 to represent the fraction of carbon in fossil fuels that is oxidized to CO₂ during the fuel combustion of petroleum, though national experts are encouraged to improve upon this assumption if better data is available. As a result, carbon mass balances for light-duty gasoline cars and trucks have been analyzed to assess what would be the most appropriate carbon oxidation fraction for these vehicles. The analysis, currently under peer review, suggests that the amount of unoxidized carbon is insignificant compared to the gaseous carbon fraction, and that 1.00 should be used to represent the oxidized carbon fraction in future inventories for gasoline fueled light-duty vehicles. Upon further peer review, the revised factor is expected to be used in future inventories. A further examination into diesel fueled vehicles is also planned.
- The 0.99 oxidation factor for coal will be further investigated in order to verify or revise this value.
- Efforts will be taken to work with EIA and other agencies to improve the quality of the U.S. territories data.

These improvements are not all-inclusive, but are part of an ongoing analysis and efforts to continually improve the CO₂ from fossil fuel combustion estimates.

3.2. Carbon Emitted from Non-Energy Uses of Fossil Fuels (IPCC Source Category 1A)

In addition to being combusted for energy, fossil fuels are also consumed for non-energy uses (NEU). These fuels are used in the industrial and transportation end-use sectors and are quite diverse, including natural gas, liquid petroleum gases (LPG), asphalt (a viscous liquid mixture of heavy crude oil distillates), petroleum coke (manufactured from heavy oil), and coal coke (manufactured from coking coal). The non-energy fuel uses are equally diverse, and include application as solvents, lubricants, and waxes, or as raw materials in the manufacture of plastics, rubber, and synthetic fibers.

Carbon dioxide emissions arise from non-energy uses via several pathways. Emissions may occur during the manufacture of a product, as is the case in producing plastics or rubber from fuel-derived feedstocks. Additionally, emissions may occur during the product's lifetime, such as during solvent use. Overall, throughout the time series and across all uses, about 65 percent of the total carbon consumed for non-energy purposes is stored in products, and not released to the atmosphere; the remaining 35 percent is emitted.

There are several areas in which non-energy uses of fossil fuels relates closely to other parts of the inventory. For example, some of the NEU products release CO₂ at the end of their commercial life when they are combusted; these emissions are reported separately within this sector in the Waste Combustion source category. In addition, there is some overlap between fossil fuels consumed for non-energy uses and the fossil-derived CO₂ emissions accounted for in the Industrial Processes sector. To avoid double-counting, the "raw" non-energy fuel consumption data reported by EIA are modified to account for these overlaps, resulting in the adjusted consumption values shown in Table 3-12. There are also net exports of petrochemicals that are not completely accounted for in the EIA data, and these affect the mass of carbon in non-energy applications; the effects of these adjustments are also shown in Table 3-11.

As shown in Table 3-11, fossil fuel emissions in 2003 from the non-energy uses of fossil fuels were 118.0 Tg CO₂ Eq., which constituted 2 percent of overall fossil fuel emissions, approximately the same proportion as in 1990. In 2003, the consumption of fuels for non-energy uses (after the adjustments described above) was 5,264 TBtu, an increase of 24 percent since 1990 (see Table 3-12). About 66.1 Tg of the C (242.5 Tg CO₂ Eq.) in these fuels was stored, while the remaining 32.2 Tg C (118.0 Tg CO₂ Eq.) was emitted. The proportion of C emitted has remained about constant since 1990, at about 31 to 35 percent of total non-energy consumption (see Table 3-13).

Table 3-11: CO₂ Emissions from Non-Energy Use Fossil Fuel Consumption (Tg CO₂ Eq.)

Year	1990	1997	1998	1999	2000	2001	2002	2003
Potential Emissions	299.3	348.3	373.3	395.0	365.8	357.8	360.9	360.5
Carbon Stored	191.4	228.0	238.0	253.4	241.1	237.7	242.1	242.5
Emissions	108.0	120.3	135.4	141.6	124.7	120.1	118.8	118.0

Methodology

The first step in estimating carbon stored in products was to determine the aggregate quantity of fossil fuels consumed for non-energy uses. The carbon content of these feedstock fuels is equivalent to potential emissions, or the product of consumption and the fuel-specific carbon content values. Both the non-energy fuel consumption and carbon content data were supplied by the EIA (2003) (see Annex 2.1). Consumption of natural gas, LPG, pentanes plus, naphthas, other oils, and special naphtha were adjusted to account for net exports of these products that are not reflected in the raw data from EIA. Consumption values for industrial coking coal, petroleum coke, other oils, and natural gas in Table 3-12 and Table 3-13, have been adjusted to subtract non-energy uses that are included in the source categories of the Industrial Processes sector.³⁴

For the remaining non-energy uses, the amount of C stored was estimated by multiplying the potential emissions by a storage factor. For several fuel types—petrochemical feedstocks (natural gas for non-fertilizer uses, LPG, pentanes plus, naphthas, other oils, still gas, special naphtha, and industrial other coal), asphalt and road oil, lubricants, and waxes—U.S. data on C stocks and flows were used to develop C storage factors, calculated as the ratio of (a) the C stored by the fuel’s non-energy products to (b) the total C content of the fuel consumed. A lifecycle approach was used in the development of these factors in order to account for losses in the production process and during use. Because losses associated with municipal solid waste management are handled separately in this sector under the Waste Combustion source category, the storage factors do not account for losses at the disposal end of the life cycle. For industrial coking coal and distillate fuel oil, storage factors were taken from the IPCC *Guidelines for National Greenhouse Gas Inventories*, which in turn draws from Marland and Rotty (1984). For the remaining fuel types (petroleum coke, miscellaneous products, and other petroleum), IPCC does not provide guidance on storage factors, and assumptions were made based on the potential fate of carbon in the respective NEU products.

³⁴ These source categories include Iron and Steel Production, Ammonia Manufacture, Carbon Black Manufacture (included in Petrochemical Production), Titanium Dioxide Production, Ferroalloy Production, and Aluminum Production.

Table 3-12: Adjusted Consumption of Fossil Fuels for Non-Energy Uses (Tbtu)

Year	1990	1997	1998	1999	2000	2001	2002	2003
Industry	3,993.2	4,794.6	5,127.6	5,398.1	4,985.1	4,933.3	5,016.4	5,023.4
Industrial Coking Coal	20.5	27.6	15.4	6.3	14.8	23.7	7.0	3.0
Industrial Other Coal	8.2	11.2	10.4	11.1	12.4	11.3	12.0	11.9
Natural Gas to Chemical Plants, Other Uses	244.1	323.7	376.3	390.3	388.9	394.0	402.0	401.0
Asphalt & Road Oil	1,170.2	1,223.6	1,262.6	1,324.4	1,275.7	1,256.9	1,240.0	1,217.4
LPG	1,001.9	1,440.9	1,568.1	1,651.2	1,497.2	1,483.9	1,550.4	1,478.4
Lubricants	186.3	182.3	190.8	192.8	189.9	174.0	171.9	160.8
Pentanes Plus	69.9	260.9	185.2	239.0	214.7	193.2	164.6	162.4
Naphtha (<401 deg. F)	294.3	467.8	529.1	458.6	556.5	473.2	559.5	588.4
Other Oil (>401 deg. F)	615.4	608.8	593.8	619.5	515.5	506.1	460.9	530.0
Still Gas	21.3	2.1	0.0	16.1	12.6	35.8	57.8	133.0
Petroleum Coke	90.7	29.9	124.7	193.9	53.9	132.8	113.6	90.9
Special Naphtha	92.1	63.8	98.1	133.9	89.1	75.5	98.7	77.5
Distillate Fuel Oil	7.0	10.4	11.7	11.7	11.7	11.7	11.7	11.7
Waxes	33.3	43.7	42.4	37.4	33.1	36.3	32.2	31.0
Miscellaneous Products	137.9	97.8	119.0	111.9	119.3	124.9	134.2	126.0
Transportation	176.0	172.1	180.2	182.1	179.4	164.3	162.4	151.8
Lubricants	176.0	172.1	180.2	182.1	179.4	164.3	162.4	151.8
U.S. Territories	86.7	92.5	94.8	114.5	165.5	80.3	80.6	88.7
Lubricants	0.7	2.5	1.3	1.4	16.4	0.0	0.0	0.0
Other Petroleum (Misc. Prod.)	86.0	90.0	93.5	113.1	149.1	80.3	80.5	88.7
Total	4,255.9	5,059.3	5,402.5	5,694.6	5,330.0	5,177.9	5,259.4	5,263.9

Note: To avoid double-counting, coal coke, petroleum coke, natural gas consumption, and other oils are adjusted for industrial process consumption reported in the Industrial Processes sector. Natural gas, LPG, Pentanes Plus, Naphthas, Special Naphtha, and Other Oils are adjusted to account for exports of chemical intermediates derived from these fuels. For residual oil (not shown in the table), all non-energy use is assumed to be consumed in carbon black production, which is also reported in the Industrial Processes sector.

- Not applicable.

Note: Totals may not sum due to independent rounding.

Table 3-13: 2003 Adjusted Non-Energy Use Fossil Fuel Consumption, Storage, and Emissions

Sector/Fuel Type	Adjusted Consumption (Tbtu)	Carbon Content (Tg C)	Storage Factor	Carbon Stored (Tg C)	Carbon Emissions (Tg C)	Carbon Emissions (Tg CO ₂ Eq.)
Industry	5,023.0	93.5	-	65.7	27.8	101.8
Industrial Coking Coal	3.0	0.1	0.75	0.1	0.0	0.1
Industrial Other Coal	11.9	0.3	0.65	0.2	0.1	0.4
Natural Gas to Chemical Plants	401.0	5.8	0.65	3.8	2.0	7.4
Asphalt & Road Oil	1,217.4	25.1	1.00	25.1	0.0	0.0
LPG	1,478.4	24.9	0.65	16.2	8.7	31.9
Lubricants	160.8	3.3	0.09	0.3	3.0	10.8
Pentanes Plus	162.4	3.0	0.65	1.9	1.0	3.8
Naphtha (<401 deg. F)	588.4	10.7	0.65	6.9	3.7	13.7
Other Oil (>401 deg. F)	530.0	10.6	0.65	6.9	3.7	13.5
Still Gas	133.0	2.3	0.65	1.5	0.8	3.0
Petroleum Coke	90.9	2.5	0.50	1.3	1.3	4.6
Special Naphtha	77.5	1.5	0.65	1.0	0.5	2.0
Distillate Fuel Oil	11.7	0.2	0.50	0.1	0.1	0.4
Waxes	31.0	0.6	0.58	0.4	0.3	0.9
Miscellaneous Products	126.0	2.5	0.00	0.0	2.5	9.3
Transportation	151.8	3.1	-	0.3	2.8	10.2
Lubricants	151.8	3.1	0.09	0.3	2.8	10.2

U.S. Territories	88.7	1.8	-	0.2	1.6	5.9
Lubricants	0.0	0.0	0.09	0.0	0.0	0.0
Other Petroleum (Misc. Prod.)	88.7	1.8	0.10	0.2	1.60	5.9
Total	5,263.9	98.3		66.1	32.2	118.0

^a To avoid double-counting, coal coke, petroleum coke, natural gas consumption, and other oils are adjusted for industrial process consumption reported in the Industrial Processes sector. Natural gas, LPG, Pentanes Plus, Naphthas, Special Naphtha, and Other Oils are adjusted to account for exports of chemical intermediates derived from these fuels. For residual oil (not shown in the table), all non-energy use is assumed to be consumed in carbon black production, which is also reported in the Industrial Processes sector.

- Not applicable.

Note: Totals may not sum due to independent rounding.

Lastly, emissions were estimated by subtracting the carbon stored from the potential emissions (see Table 3-11). More detail on the methodology for calculating storage and emissions from each of these sources is provided in Annex 2.3.

Where storage factors were calculated specifically for the United States, data were obtained on fuel products such as asphalt, plastics, synthetic rubber, synthetic fibers, carbon black, personal cleansers, pesticides, and solvents, and industrial releases including VOC, solvent, and non-combustion CO emissions, TRI releases, refinery wastewater, hazardous waste incineration, and energy recovery. Data were taken from a variety of industry sources, government reports, and expert communications. Sources include EPA's compilations of air emission factors (EPA 1995, 2001), *National Air Quality and Emissions Trends Report* data (EPA 2004a), *Toxics Release Inventory, 1998* (2000a), *Biennial Reporting System* data (EPA 2004b), pesticide sales and use estimates (EPA 1998, 1999, 2002) and hazardous waste data (EPA 2004b); the EIA Manufacturer's Energy Consumption Survey (MECS) (EIA 1994, 1997, 2001b); the National Petrochemical & Refiners Association (NPRA 2001); the National Asphalt Pavement Association (Connolly 2000); the Emissions Inventory Improvement Program (EIIP 1998, 1999); the U.S. Bureau of the Census (1999, 2003); the American Plastics Council (APC 2000, 2001, 2003; Eldredge-Roebuck 2000); the Society of the Plastics Industry (SPI 2000); the Rubber Manufacturers' Association (RMA 2002; STMC 2003); the International Institute of Synthetic Rubber Products (IISRP 2000); the Fiber Economics Bureau (FEB 2001); the; *Material Safety Data Sheets* (Miller 1999); and the Chemical Manufacturer's Association (CMA 1999); Specific data sources are listed in full detail in Annex 2.3.

Uncertainty

A Tier 2 Monte Carlo analysis was performed using @RISK software to determine the level of uncertainty surrounding the estimates of emissions and storage factors from non-energy uses. The Tier 2 analysis was performed to allow the specification of probability density functions for key variables, within a computational structure that mirrors the calculation of the inventory estimate.

As noted above, the non-energy use analysis is based on U.S.-specific storage factors for (1) feedstock materials (natural gas, LPG, pentanes plus, naphthas, other oils, still gas, special naphthas, and other industrial coal), (2) asphalt, (3) lubricants, and (4) waxes. For the remaining fuel types (the "other" category), the storage factors were taken directly from the IPCC *Guidelines for National Greenhouse Gas Inventories*, where available, and otherwise assumptions were made based on the potential fate of carbon in the respective NEU products. To characterize uncertainty, five separate analyses were conducted, corresponding to each of the five categories. In all cases, statistical analyses or expert judgments of uncertainty were not available directly from the information sources for all the activity variables; thus, uncertainty estimates were determined using assumptions based on source category knowledge.

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-14 (emissions) and Table 3-15 (storage factors). Carbon emitted from non-energy uses of fossil fuels in 2003 was estimated to be between 97.5 and 130.9 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 17 percent below to 11 percent above the 2003 emission estimate of 118.0 Tg CO₂ Eq. The uncertainty in the emission estimates are a function of uncertainty in both the quantity of fuel used for non-energy purposes and the storage factor.

Table 3-14: Tier 2 Quantitative Uncertainty Estimates for CO₂ Emissions from Non-Energy Uses of Fossil Fuels (Tg CO₂ Eq. and Percent)

Source	Gas	2003 Emission Estimate (Tg CO ₂ Eq.)	Uncertainty Range Relative to Emission Estimate ^a			
			(Tg CO ₂ Eq.)		(%)	
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Feedstocks	CO ₂	75.3	61.0	90.5	-19%	+20%
Asphalt	CO ₂	0.0	0.2	0.9	NA	NA
Lubricants	CO ₂	21.1	17.4	24.4	-17%	+16%
Waxes	CO ₂	0.9	0.7	1.5	-24%	+54%
Other	CO ₂	20.9	8.9	23.2	-57%	+11%
Total	CO₂	118.0	97.5	130.9	-17%	+11%

^aRange of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

NA (Not Applicable)

Table 3-15: Tier 2 Quantitative Uncertainty Estimates for Storage Factors of Non-Energy Uses of Fossil Fuels (Percent)

Source	Gas	2003 Storage Factor (%)	Uncertainty Range Relative to Inventory Factor ^a			
			(%)		(% , Relative)	
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Feedstocks	CO ₂	65%	63%	67%	-3%	+3%
Asphalt	CO ₂	100%	99%	100%	-1%	+0%
Lubricants	CO ₂	9%	4%	18%	-58%	+90%
Waxes	CO ₂	58%	44%	69%	-24%	+19%
Other	CO ₂	24%	18%	67%	-24%	+180%
Total	CO₂	65%	63%	67%	-3%	+3%

^aRange of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

In Table 3-15, feedstocks and asphalt contribute least to overall storage factor uncertainty. Although the feedstocks category—the largest use category in terms of total carbon flows—appears to have tight confidence limits, this is to some extent an artifact of the way the uncertainty analysis was structured. As discussed in Annex 2.3, the storage factor for feedstocks is based on an analysis of five fates that result in long-term storage (e.g., plastics production), and ten that result in emissions (e.g., volatile organic compound emissions). Rather than modeling the total uncertainty around all 15 of these fate processes, the current analysis addresses only the storage fates, and assumes that all C that is not stored is emitted. As the production statistics that drive the storage values are relatively well-characterized, this approach yields a result that is probably biased toward understating uncertainty.

As is the case with the other uncertainty analyses discussed throughout this document, the uncertainty results above address only those factors that can be readily quantified. More details on the uncertainty analysis are provided in Annex 2.3.

QA/QC and Verification

A source-specific QA/QC plan for non-energy uses of fossil fuels was developed and implemented. This effort included a Tier 1 analysis, as well as portions of a Tier 2 analysis for non-energy uses involving petrochemical feedstocks. The Tier 2 procedures that were implemented involved checks specifically focusing on the activity data and methodology for estimating the fate of C (in terms of storage and emissions) across the various end-uses of fossil carbon. Emission and storage totals for the different subcategories were compared, and trends across the time series were carefully analyzed to determine whether any corrective actions were needed. Corrective actions were taken to rectify minor errors and to improve the transparency of the calculations, facilitating future QA/QC.

Recalculations Discussion

This year's methodology reflects several refinements and improvements. First and most fundamentally, this year the NEU analysis is presented as its own source category. In the past, the NEU component of this category was described in the context of a calculation sequence that first determined potential emissions (based on total fuel consumption for all purposes), and then deducted (1) C not oxidized (e.g., in ash) and (2) C put into long-term storage as a result of NEUs. In this context, the focus of the NEU section was previously on carbon storage rather than emissions. This year, the presentation within this source category emphasizes emissions, though the storage factors (i.e., fraction of C stored rather than emitted) are still presented to facilitate comparison with IPCC guidance.

Several substantive changes also appear in this year's inventory. First, a U.S.-specific storage factor (58 percent) was developed for waxes. Previously, the IPCC storage factor for wax (100 percent stored) had been used.

In addition, the methodology for calculating emissions and storage for feedstocks has been revised in several ways. First, three additional fuel types have been added as inputs to the system covered by the feedstocks mass balance calculations: other industrial coal, still gas, and special naphthas. The other significant improvement for feedstocks is that the estimates of U.S. plastics consumption have been revised: in previous years, the consumption data for some of the plastic resins in the dataset included consumption in Canada and Mexico. By adjusting the data to reflect U.S. (rather than North American) consumption for those resins, the accuracy of the estimate for feedstocks has improved. In addition, several minor adjustments were made (e.g., the calculations for storage now include synthetic rubber in durable and non-durable goods other than tires, and the calculations for emissions include losses due to abrasion/ oxidation of rubber from tires).

As noted in the beginning of the section, there are several areas where the boundaries of the NEU analysis adjoin the systems covered within the Industrial Processes sector. Carbon black production has been removed from the NEU analysis, as it is now addressed as a subcategory of IP-petrochemical emissions (if it was retained in NEU, there would be double-counting). As a result of this change, residual oil—which is an input to carbon black production, and had been among the fuels covered in the NEU section in previous years—no longer appears in the list of fuels in this section. A portion of other oils is also assumed to be used in the production of carbon black, and therefore other oils consumption is adjusted to account for the amount of fuel used in the C black calculation.

The storage factor for miscellaneous products also changed. In prior years, a value of 100 percent was used; in this year's inventory, based on a review of the specific applications of these products, there is now assumed to be no storage at all from these products.

There have been several updates to the data used to calculate storage factors, not only by adding information for 2003 (where available) but also for updating data sets for earlier years. For example, the results reflect new data for hazardous waste incineration (1999 through 2001) and fiber production (2001 through 2002).

Planned Improvements

There are several improvements planned for the future:

- Collecting additional information on energy recovery from Manufacturing Energy Consumption Surveys. An effort is planned to carefully examine the “microdata” from these surveys to determine the nature and quantity of materials initially identified as being destined for “non-energy use” that are actually combusted for energy recovery.
- Improving the uncertainty analysis. Most of the input parameter distributions are based on professional judgment rather than rigorous statistical characterizations of uncertainty.
- Better characterizing flows of fossil carbon. Additional “fates” may be researched, including: the fossil carbon load in organic chemical wastewaters; an expanded import and export analysis (i.e., evaluating additional commodities); and improving the characterization of cleansers (to exclude any potential biogenic carbon sources).

Finally, although U.S.-specific storage factors have been developed for feedstocks, asphalt, lubricants, and waxes, default values from IPCC are still used for two of the non-energy fuel types (industrial coking coal and distillate oil), and broad assumptions are being used for the remaining fuels (petroleum coke, miscellaneous products, and other petroleum). Over the long term, there are plans to improve these storage factors by conducting analyses of C fate similar to those described in Annex 2.3.

3.3. Stationary Combustion (excluding CO₂) (IPCC Source Category 1A)

Stationary combustion encompasses all fuel combustion activities from fixed sources (versus mobile combustion). Other than CO₂, which was addressed in the previous section, gases from stationary combustion include the greenhouse gases CH₄ and N₂O and the ambient air pollutants NO_x, CO, and NMVOCs.³⁵ Emissions of these gases from stationary combustion sources depend upon fuel characteristics, size and vintage, along with combustion technology, pollution control equipment, and ambient environmental conditions. Emissions also vary with operation and maintenance practices.

Nitrous oxide and NO_x emissions from stationary combustion are closely related to air-fuel mixes and combustion temperatures, as well as the characteristics of any pollution control equipment that is employed. Carbon monoxide emissions from stationary combustion are generally a function of the efficiency of combustion; they are highest when less oxygen is present in the air-fuel mixture than is necessary for complete combustion. These conditions are most likely to occur during start-up, shutdown and during fuel switching (e.g., the switching of coal grades at a coal-burning electric utility plant). Methane and NMVOC emissions from stationary combustion are primarily a function of the CH₄ and NMVOC content of the fuel and combustion efficiency.

Emissions of CH₄ decreased 15 percent overall to 6.7 Tg CO₂ Eq. (319 Gg) in 2003. This decrease in CH₄ emissions was primarily due to lower wood consumption in the residential sector. Conversely, N₂O emissions rose 13 percent since 1990 to 13.8 Tg CO₂ Eq. (45 Gg) in 2003. The largest source of N₂O emissions was coal combustion by electricity generators, which alone accounted for 63 percent of total N₂O emissions from stationary combustion in 2003. Overall, however, stationary combustion is a small source of CH₄ and N₂O in the United States.

In contrast, stationary combustion is a significant source of NO_x emissions, though a smaller source of CO and NMVOCs. In 2003, emissions of NO_x from stationary combustion represented 39 percent of national NO_x emissions, while CO and NMVOC emissions from stationary combustion contributed approximately 5 and 7 percent, respectively, to the national totals. From 1990 to 2003, emissions of NO_x and CO from stationary combustion decreased by 27 and 11 percent, respectively, and emissions of NMVOCs increased by 10 percent.

The decrease in NO_x emissions from 1990 to 2003 are mainly due to decreased emissions from electric power. The decrease in CO and increase in NMVOC emissions over this time period can largely be attributed to apparent changes in residential wood use, which is the most significant source of these pollutants from stationary combustion. Table 3-16 through Table 3-19 provide CH₄ and N₂O emission estimates from stationary combustion by sector and fuel type. Estimates of NO_x, CO, and NMVOC emissions in 2003 are given in Table 3-20.³⁶

Table 3-16: CH₄ Emissions from Stationary Combustion (Tg CO₂ Eq.)

Sector/Fuel Type	1990	1997	1998	1999	2000	2001	2002	2003
Electric Power	0.6	0.6	0.7	0.7	0.7	0.7	0.7	0.7
Coal	0.3	0.4	0.4	0.4	0.4	0.4	0.4	0.4
Fuel Oil	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Natural gas	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1

³⁵ Sulfur dioxide (SO₂) emissions from stationary combustion are addressed in Annex 6.3.

³⁶ See Annex 3.1 for a complete time series of ambient air pollutant emission estimates for 1990 through 2003.

Wood	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Industrial	2.1	2.4	2.3	2.2	2.3	2.1	2.1	2.1
Coal	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Fuel Oil	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Natural gas	0.8	0.9	0.9	0.9	0.9	0.8	0.8	0.8
Wood	0.9	1.0	1.0	1.0	1.0	0.9	0.9	0.9
Commercial	0.7	0.8	0.8	0.8	0.8	0.7	0.7	0.8
Coal	+	+	+	+	+	+	+	+
Fuel Oil	0.2	0.1	0.1	0.1	0.2	0.2	0.2	0.2
Natural gas	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Wood	0.2	0.3	0.3	0.3	0.3	0.2	0.3	0.3
Residential	4.4	3.5	3.1	3.4	3.5	3.1	2.8	3.1
Coal	0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Fuel Oil	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Natural Gas	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Wood	3.5	2.6	2.3	2.5	2.6	2.2	1.9	2.1
U.S. Territories	+	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Coal	+	+	+	+	+	+	+	+
Fuel Oil	+	0.1	0.1	0.1	+	0.1	0.1	0.1
Natural Gas	+	+	+	+	+	+	+	+
Wood	+	+	+	+	+	+	+	+
Total	7.8	7.4	6.9	7.1	7.3	6.7	6.4	6.7

+ Does not exceed 0.05 Tg CO₂ Eq.

Note: Totals may not sum due to independent rounding.

Table 3-17: N₂O Emissions from Stationary Combustion (Tg CO₂ Eq.)

Sector/Fuel Type	1990	1997	1998	1999	2000	2001	2002	2003
Electric Power	7.6	8.6	8.9	8.9	9.3	9.0	9.1	9.3
Coal	7.1	8.2	8.3	8.3	8.7	8.5	8.6	8.7
Fuel Oil	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2
Natural Gas	0.1	0.1	0.1	0.1	0.2	0.2	0.2	0.2
Wood	0.2	0.2	0.2	0.2	0.2	0.1	0.2	0.2
Industrial	3.2	3.5	3.3	3.3	3.3	3.1	3.2	3.2
Coal	0.7	0.7	0.7	0.6	0.6	0.6	0.6	0.6
Fuel Oil	0.5	0.5	0.5	0.5	0.5	0.6	0.6	0.6
Natural Gas	0.2	0.3	0.3	0.3	0.3	0.2	0.2	0.2
Wood	1.7	2.0	1.9	1.9	1.9	1.7	1.8	1.8
Commercial	0.4	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Coal	0.1	0.1	+	+	+	+	+	+
Fuel Oil	0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Natural Gas	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Wood	+	0.1	0.1	0.1	0.1	+	+	+
Residential	1.1	0.9	0.8	0.9	1.0	0.9	0.8	0.9
Coal	+	+	+	+	+	+	+	+
Fuel Oil	0.3	0.3	0.2	0.3	0.3	0.3	0.3	0.3
Natural Gas	0.1	0.2	0.1	0.1	0.2	0.1	0.1	0.2
Wood	0.7	0.5	0.5	0.5	0.5	0.4	0.4	0.4
U.S. Territories	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Coal	+	+	+	+	+	+	+	+
Fuel Oil	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Natural Gas	+	+	+	+	+	+	+	+
Wood	+	+	+	+	+	+	+	+
Total	12.3	13.5	13.4	13.5	14.0	13.5	13.5	13.8

+ Does not exceed 0.05 Tg CO₂ Eq.

Note: Totals may not sum due to independent rounding.

Table 3-18: CH₄ Emissions from Stationary Combustion (Gg)

Sector/Fuel Type	1990	1997	1998	1999	2000	2001	2002	2003
Electric Power	27	29	31	31	32	32	32	33
Coal	16	19	19	19	20	19	20	20
Fuel Oil	4	3	4	4	3	4	3	4
Natural Gas	3	4	4	5	5	5	5	5
Wood	4	4	4	4	4	4	4	5
Industrial	101	115	108	107	108	100	102	100
Coal	17	16	15	14	15	14	13	13
Fuel Oil	6	6	5	5	6	6	6	6
Natural Gas	37	44	43	41	42	38	39	37
Wood	41	49	46	46	47	41	44	43
Commercial	35	38	36	37	39	35	35	36
Coal	1	1	1	1	1	1	1	1
Fuel Oil	10	7	7	7	8	7	7	8
Natural Gas	13	16	15	15	16	15	15	15
Wood	11	14	14	15	15	12	12	12
Residential	209	167	150	160	167	148	132	146
Coal	8	5	4	4	4	4	3	4
Fuel Oil	14	14	13	15	16	15	15	15
Natural Gas	21	24	22	23	24	23	24	25
Wood	166	123	110	118	123	105	89	102
U.S. Territories	2	2	2	2	2	3	3	4
Coal	+	+	+	+	+	+	+	+
Fuel Oil	2	2	2	2	2	3	3	3
Natural Gas	+	+	+	+	+	+	+	+
Wood	+	+	+	+	+	+	+	+
Total	373	351	328	338	349	318	305	319

+ Does not exceed 0.5 Gg

Note: Totals may not sum due to independent rounding.

Table 3-19: N₂O Emissions from Stationary Combustion (Gg)

Sector/Fuel Type	1990	1997	1998	1999	2000	2001	2002	2003
Electricity Generation	24	28	29	29	30	29	29	30
Coal	23	26	27	27	28	27	28	28
Fuel Oil	1	1	1	1	1	1	1	1
Natural Gas	+	+	+	+	1	1	1	+
Wood	+	1	1	1	1	+	1	1
Industrial	10	11	11	11	11	10	10	10
Coal	2	2	2	2	2	2	2	2
Fuel Oil	2	2	2	2	2	2	2	2
Natural Gas	1	1	1	1	1	1	1	1
Wood	5	7	6	6	6	5	6	6
Commercial	1	1	1	1	1	1	1	1
Coal	+	+	+	+	+	+	+	+
Fuel Oil	1	+	+	+	+	+	+	+
Natural Gas	+	+	+	+	+	+	+	+
Wood	+	+	+	+	+	+	+	+
Residential	4	3	3	3	3	3	3	3
Coal	+	+	+	+	+	+	+	+
Fuel Oil	1	1	1	1	1	1	1	1
Natural Gas	+	+	+	+	+	+	+	+

Wood	2		2	1	2	2	1	1	1
U.S. Territories	+		+	+	+	+	+	+	+
Coal	+		+	+	+	+	+	+	+
Fuel Oil	+		+	+	+	+	+	+	+
Natural Gas	+		+	+	+	+	+	+	+
Wood	+		+	+	+	+	+	+	+
Total	40		44	43	43	45	43	44	45

+ Does not exceed 0.5 Gg

Note: Totals may not sum due to independent rounding.

Table 3-20: NO_x, CO, and NMVOC Emissions from Stationary Combustion in 2003 (Gg)

Sector/Fuel Type	NO_x	CO	NMVOC
Electric Generation	4,045	480	50
Coal	3,447	240	24
Fuel Oil	135	30	4
Natural gas	301	101	11
Wood	34	NA	NA
Other Fuels ^a	NA	35	2
Internal Combustion	127	73	10
Industrial	2,516	1,249	154
Coal	522	138	10
Fuel Oil	154	50	8
Natural gas	920	402	52
Wood	NA	NA	NA
Other Fuels ^a	116	353	28
Internal Combustion	803	307	55
Commercial/Institutional	244	149	29
Coal	19	13	1
Fuel Oil	49	17	3
Natural gas	155	82	13
Wood	NA	NA	NA
Other Fuels ^a	21	37	11
Residential	417	2,575	773
Coal ^b	NA	NA	NA
Fuel Oil ^b	NA	NA	NA
Natural Gas ^b	NA	NA	NA
Wood	20	2,358	748
Other Fuels	397	217	25
Total	7,222	4,454	1,007

NA (Not Available)

^a Includes LPG, waste oil, coke oven gas, and coke (EPA 2003), (EPA 2004a).

^b Residential coal, fuel oil, and natural gas emissions are included in "Other Fuels" (EPA 2003), (EPA 2004a).

Note: Totals may not sum due to independent rounding. See Annex 3.1 for emissions in 1990 through 2003.

Methodology

Methane and N₂O emissions were estimated by multiplying fossil fuel and wood consumption data by emission factors (by sector and fuel type). National coal, natural gas, fuel oil, and wood consumption data were grouped by sector: industrial, commercial, residential, electric power, and U.S. territories. For the CH₄ and N₂O estimates, fuel consumption data for the United States were obtained from EIA's *Monthly Energy Review* and unpublished

supplemental tables on petroleum product detail (EIA 2004). Because the United States does not include territories in its national energy statistics, fuel consumption data for territories were provided separately by the Grillo (2004).³⁷ Fuel consumption for the industrial sector was adjusted to subtract out construction and agricultural use, which is reported under mobile sources.³⁸ Construction and agricultural fuel use was obtained from EPA (2004b). Estimates for wood biomass consumption for fuel combustion do not include wood wastes, liquors, municipal solid waste, tires, etc. that are reported as biomass by EIA.

Emission factors for the four end-use sectors were provided by the *Revised 1996 IPCC Guidelines for National Greenhouse Gas Inventories* (IPCC/UNEP/OECD/IEA 1997). U.S. territories' emission factors were estimated using the U.S. emission factors for the primary sector in which each fuel was combusted.

Emission estimates for NO_x, CO, and NMVOCs in this section were obtained from preliminary data (EPA 2004a) and disaggregated based on EPA (2003), which, in its final iteration, will be published on the National Emission Inventory (NEI) Air Pollutant Emission Trends web site. The major categories included in this section are those reported in EPA (2003) and EPA (2004a): coal, fuel oil, natural gas, wood, other fuels (including LPG, coke, coke oven gas, and others), and stationary internal combustion. The EPA estimates emissions of NO_x, CO, and NMVOCs by sector and fuel source using a "bottom-up" estimating procedure. In other words, emissions were calculated either for individual sources (e.g., industrial boilers) or for multiple sources combined, using basic activity data as indicators of emissions. Depending on the source category, these basic activity data may include fuel consumption, fuel deliveries, tons of refuse burned, raw material processed, etc.

The overall emission control efficiency of a source category was derived from published reports, the 1985 National Acid Precipitation and Assessment Program (NAPAP) emissions inventory, and other EPA databases. The U.S. approach for estimating emissions of NO_x, CO, and NMVOCs from stationary combustion, as described above, is consistent with the methodology recommended by the IPCC (IPCC/UNEP/OECD/IEA 1997).

More detailed information on the methodology for calculating emissions from stationary combustion, including emission factors and activity data, is provided in Annex 3.1.

Uncertainty

Methane emission estimates from stationary sources exhibit high uncertainty, primarily due to difficulties in calculating emissions from wood combustion (i.e., fireplaces and wood stoves). The estimates of CH₄ and N₂O emissions presented are based on broad indicators of emissions (i.e., fuel use multiplied by an aggregate emission factor for different sectors), rather than specific emission processes (i.e., by combustion technology and type of emission control).

An uncertainty analysis was performed by primary fuel type for each end-use sector, using the IPCC-recommended Tier 2 uncertainty estimation methodology, Monte Carlo Simulation technique, with @RISK software.

The uncertainty estimation model for this source category was developed by integrating the CH₄ and N₂O stationary source inventory estimation models with the model for CO₂ from fossil fuel combustion to realistically characterize the interaction (or endogenous correlation) between the variables of these three models. A total of 115 input variables were simulated for the uncertainty analysis of this source category (85 from the CO₂ emissions from fossil fuel combustion inventory estimation model and 30 from the stationary source inventory models).

³⁷ U.S. territories data also include combustion from mobile activities because data to allocate territories' energy use were unavailable. For this reason, CH₄ and N₂O emissions from combustion by U.S. territories are only included in the stationary combustion totals.

³⁸ Though emissions from construction and farm use occur due to both stationary and mobile sources, detailed data was not available to determine the magnitude from each. Currently, these emissions are assumed to be predominantly from mobile sources.

In developing the uncertainty estimation model, uniform distribution was assumed for all activity-related input variables and N₂O emission factors, based on the SAIC/EIA (2001) report.³⁹ For these variables, the uncertainty ranges were assigned to the input variables based on the data reported in SAIC/EIA (2001).⁴⁰ However, the CH₄ emission factors differ from those used by EIA. Since these factors were obtained from IPCC/UNEP/OECD/IEA (1997), uncertainty ranges were assigned based on IPCC default uncertainty estimates (IPCC Good Practice Guidance 2000).

The uncertainty ranges for the activity-related input variables and N₂O emission factors were typically asymmetric around their inventory estimates. The uncertainty ranges for the non-biomass-related CH₄ emission factors were symmetric around their inventory estimates; for biomass, they were asymmetric around their emission factor estimates. Bias (or systematic uncertainties) associated with these variables accounted for much of the uncertainty associated with the activity data and N₂O emission factor variables.⁴¹ For purposes of this uncertainty analysis, each input variable was simulated 10,000 times through Monte Carlo sampling.

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-21. Stationary combustion CH₄ emissions in 2003 (*including* biomass) were estimated to be between 4.8 and 13.4 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 28 percent below to 99 percent above the 2003 emission estimate of 6.7 Tg CO₂ Eq.⁴² Stationary combustion N₂O emissions in 2003 (*including* biomass) were estimated to be between 10.9 and 39.5 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 22 percent below to 184 percent above the 2003 emissions estimate of 13.8 Tg CO₂ Eq.

Table 3-21: Tier 2 Quantitative Uncertainty Estimates for CH₄ and N₂O Emissions from Energy-Related Stationary Combustion, Including Biomass (Tg CO₂ Eq. and Percent)

Combustion, including Biomass (Tg CO ₂ Eq. and Percent)						
Source	Gas	2003 Emission				
		Estimate (Tg CO ₂ Eq.)	Uncertainty Range Relative to Emission Estimate ^a			
			(Tg CO ₂ Eq.)		(%)	
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Stationary Combustion	CH ₄	6.7	4.8	13.4	-28%	+99%
Stationary Combustion	N ₂ O	13.8	10.9	39.5	-22%	+184%

^a Range of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

The uncertainties associated with the emission estimates of CH₄ and N₂O are greater than those associated with estimates of CO₂ from fossil fuel combustion, which mainly rely on the carbon content of the fuel combusted. Uncertainties in both CH₄ and N₂O estimates are due to the fact that emissions are estimated based on emission factors representing only a limited subset of combustion conditions. For the ambient air pollutants, uncertainties are

³⁹ SAIC/EIA(2001) characterizes the underlying probability density function for the input variables as a combination of uniform and normal distributions (the former distribution to represent the bias component and the latter to represent the random component). However, for purposes of the current uncertainty analysis, it was determined that uniform distribution was more appropriate to characterize the probability density function underlying each of these variables.

⁴⁰ In the SAIC/EIA (2001) report, the quantitative uncertainty estimates were developed for each of the three major fossil fuels used within each end-use sector; the variations within the sub-fuel types within each end-use sector were not modeled. However, for purposes of assigning uncertainty estimates to the sub-fuel type categories within each end-use sector in the current uncertainty analysis, SAIC/EIA (2001)-reported uncertainty estimates were extrapolated.

⁴¹ Although, in general, random uncertainties are the main focus of statistical uncertainty analysis, when the uncertainty estimates are elicited from experts, their estimates include both random and systematic uncertainties. Hence, both these types of uncertainties are represented in this uncertainty analysis.

⁴² The low emission estimates reported in this section have been rounded down to the nearest integer values and the high emission estimates have been rounded up to the nearest integer values.

partly due to assumptions concerning combustion technology types, age of equipment, emission factors used, and activity data projections.

QA/QC and Verification

A source-specific QA/QC plan for stationary combustion was developed and implemented. This effort included a Tier 1 analysis, as well as portions of a Tier 2 analysis. The Tier 2 procedures that were implemented involved checks specifically focusing on the activity data and emission factor sources and methodology used for estimating CH₄, N₂O, and the ambient air pollutants from stationary combustion in the United States. Emission totals for the different sectors and fuels were compared and trends were investigated.

A few corrective actions were taken. It was determined that emissions from construction and farm use had been previously estimated in both this source category (in the industrial sector) and for mobile sources. To avoid double-counting, these emissions are now reported only under mobile sources. Therefore, it was necessary to subtract out energy consumption from construction and agriculture from the industrial sector. Also, a small error was corrected in the conversion of natural gas and wood energy consumption from gross calorific value to net calorific value.

Recalculations Discussion

Historical CH₄ and N₂O emissions from stationary sources (excluding CO₂) were revised due to several changes. First, industrial sector energy consumption was adjusted downward to avoid double-counting of emissions from construction and agriculture. Second, the conversion of natural gas and wood energy consumption from gross calorific value to net calorific value was corrected. These adjustments have been explained in the previous section. Third, slight changes to emission estimates for the other sectors are due to revised data from EIA (2004). This latter revision is explained in greater detail in the section on CO₂ Emissions from Fossil Fuel Combustion within this sector. The combination of the methodological and historical data changes resulted in an average annual decrease of 0.4 Tg CO₂ Eq. (5.2 percent) in CH₄ emissions from stationary combustion and an average annual decrease of 0.4 Tg CO₂ Eq. (3.2 percent) in N₂O emissions from stationary combustion for the period 1990 through 2002.

Planned Improvements

Several items are being evaluated to improve the CH₄ and N₂O emission estimates from stationary source combustion and to reduce uncertainty. Efforts will be taken to work with EIA and other agencies to improve the quality of the U.S. territories data. Because these data are not broken out by stationary and mobile uses, further research will be aimed at trying to allocate consumption appropriately. In addition, the uncertainty of biomass emissions will be further investigated. Currently, the exclusion of biomass increases the uncertainty, although it was expected to reduce the uncertainty. These improvements are not all-inclusive, but are part of an ongoing analysis and efforts to continually improve these stationary estimates.

3.4. Mobile Combustion (excluding CO₂) (IPCC Source Category 1A)

Mobile combustion emits greenhouse gases other than CO₂, including CH₄, N₂O, and the ambient air pollutants NO_x, CO, and NMVOCs. While air conditioners and refrigerated units in vehicles also emit hydrofluorocarbons (HFCs), these gases are covered in Chapter 3, Industrial Processes, under the section entitled Substitution of Ozone Depleting Substances. As with stationary combustion, N₂O and NO_x emissions are closely related to fuel characteristics, air-fuel mixes, combustion temperatures, as well as usage of pollution control equipment. Nitrous oxide, in particular, can be formed by the catalytic processes used to control NO_x, CO, and hydrocarbon emissions. Carbon monoxide emissions from mobile combustion are significantly affected by combustion efficiency and the presence of post-combustion emission controls. Carbon monoxide emissions are highest when air-fuel mixtures have less oxygen than required for complete combustion. These emissions occur especially in idle, low speed, and cold start conditions. Methane and NMVOC emissions from motor vehicles are a function of the CH₄ content of the motor fuel, the amount of hydrocarbons passing uncombusted through the engine, and any post-combustion control of hydrocarbon emissions, such as catalytic converters.

Emissions from mobile combustion were estimated by transport mode (e.g., highway, air, rail), fuel type (e.g. motor gasoline, diesel fuel, jet fuel), and vehicle type (e.g. passenger cars, light-duty trucks). Road transport accounted for the majority of mobile source fuel consumption, and hence, the majority of mobile combustion emissions. Table 3-22 and Table 3-23 provide CH₄ and N₂O emission estimates, respectively, in Tg CO₂ Eq.; Table 3-24 and Table 3-25 present these estimates in Gg of each gas. Estimates of NO_x, CO, and NMVOC emissions are given in Table 3-26 through Table 3-28.⁴³

Mobile combustion was responsible for a small portion of national CH₄ emissions (0.5 percent) but was the second largest source of N₂O (11.2 percent) in the United States. From 1990 to 2003, CH₄ emissions declined by 44 percent, to 2.7 Tg CO₂ Eq. (128 Gg), due largely to control technologies employed on highway vehicles in the United States that reduce CO, NO_x, NMVOC, and CH₄ emissions. The same technologies, however, resulted in higher N₂O emissions, causing a 27 percent increase in N₂O emissions from mobile sources between 1990 and 1998. Nitrous oxide emissions have subsequently declined 24 percent as improvements in the emission control technologies installed on new vehicles have reduced emission rates of both NO_x and N₂O per vehicle mile traveled. As a result, N₂O emissions in 2003 were 4 percent lower than in 1990, at 42.1 Tg CO₂ Eq. (136 Gg) (see Figure 3-17). Overall, CH₄ and N₂O emissions were predominantly from gasoline-fueled passenger cars and light-duty gasoline trucks.

Figure 3-17: Mobile Source CH₄ and N₂O Emissions

Table 3-22: CH₄ Emissions from Mobile Combustion (Tg CO₂ Eq.)

Fuel Type/Vehicle Type ^a	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	4.3	3.5	3.3	3.0	2.7	2.4	2.2	2.0
Passenger Cars	2.6	1.9	1.8	1.7	1.5	1.4	1.2	1.1
Light-Duty Trucks	1.4	1.3	1.3	1.1	1.0	0.9	0.8	0.8
Heavy-Duty Vehicles	0.3	0.2	0.2	0.2	0.1	0.1	0.1	0.1
Motorcycles	+	+	+	+	+	+	+	+
Diesel Highway	+	+	+	+	+	+	+	+
Passenger Cars	+	+	+	+	+	+	+	+
Light-Duty Trucks	+	+	+	+	+	+	+	+
Heavy-Duty Vehicles	+	+	+	+	+	+	+	+
Alternative Fuel Highway	+	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Non-Highway	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Ships and Boats	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Locomotives	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Farm Equipment	0.2	0.2	0.1	0.2	0.2	0.1	0.1	0.1
Construction Equipment	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Aircraft	+	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Other ^b	+	+	+	+	+	+	0.1	0.1
Total	4.8	4.0	3.9	3.6	3.4	3.1	2.9	2.7

+ Less than 0.05 Tg CO₂ Eq.

Note: Totals may not sum due to independent rounding.

^a See Annex 3.2 for definitions of highway vehicle types.

^b "Other" includes snowmobiles and other recreational equipment, logging equipment, lawn and garden equipment, railroad equipment, airport equipment, commercial equipment, and industrial equipment.

Table 3-23: N₂O Emissions from Mobile Combustion (Tg CO₂ Eq.)

Fuel Type/Vehicle Type	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	40.3	51.5	51.7	50.8	49.1	45.0	41.6	38.0

⁴³ See Annex 3.2 for a complete time series of emission estimates for 1990 through 2003.

Passenger Cars	25.5		26.7	26.7	25.9	24.7	23.1	21.6	19.8
Light-Duty Trucks	14.1		23.7	23.7	23.6	23.0	20.6	18.6	16.7
Heavy-Duty Vehicles	0.7		1.1	1.3	1.3	1.3	1.4	1.4	1.4
Motorcycles	+		+	+	+	+	+	+	+
Diesel Highway	0.2		0.3	0.3	0.3	0.3	0.3	0.3	0.3
Passenger Cars	+		+	+	+	+	+	+	+
Light-Duty Trucks	+		+	+	+	+	+	+	+
Heavy-Duty Vehicles	0.2		0.2	0.3	0.3	0.3	0.3	0.3	0.3
Alternative Fuel Highway	+		0.1	0.1	0.1	0.1	0.1	0.1	0.1
Non-Highway	3.2		3.3	3.3	3.4	3.7	3.5	3.6	3.6
Ships and Boats	0.4		0.3	0.3	0.3	0.5	0.3	0.5	0.5
Locomotives	0.3		0.3	0.3	0.3	0.3	0.3	0.3	0.3
Farm Equipment	1.7		1.7	1.8	1.8	1.9	1.8	1.7	1.7
Construction Equipment	0.2		0.3	0.3	0.3	0.3	0.3	0.3	0.3
Aircraft	0.3		0.4	0.4	0.4	0.4	0.4	0.4	0.5
Other*	0.2		0.3	0.3	0.3	0.3	0.3	0.3	0.3
Total	43.7		55.2	55.3	54.6	53.2	49.0	45.6	42.1

+ Less than 0.05 Tg CO₂ Eq.

Note: Totals may not sum due to independent rounding.

*"Other" includes snowmobiles and other recreational equipment, logging equipment, lawn and garden equipment, railroad equipment, airport equipment, commercial equipment, and industrial equipment.

Table 3-24: CH₄ Emissions from Mobile Combustion (Gg)

Fuel Type/Vehicle Type	1990		1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	205		166	158	143	130	116	105	95
Passenger Cars	125		91	87	82	73	65	59	53
Light-Duty Trucks	65		64	60	52	49	44	40	38
Heavy-Duty Vehicles	13		10	10	9	7	6	5	5
Motorcycles	1		1	1	1	1	1	1	1
Diesel Highway	1		1	1	1	1	1	1	1
Passenger Cars	+		+	+	+	+	+	+	+
Light-Duty Trucks	+		+	+	+	+	+	+	+
Heavy-Duty Vehicles	1		1	1	1	1	1	1	1
Alternative Fuel Highway	1		3	3	4	4	5	5	6
Non-Highway	22		23	23	24	26	25	26	26
Ships and Boats	4		3	3	3	5	3	4	4
Locomotives	3		3	3	3	3	3	3	3
Farm Equipment	7		7	7	7	7	7	7	7
Construction Equipment	4		5	5	5	5	6	6	6
Aircraft	2		3	3	3	3	3	3	3
Other*	2		2	2	2	2	2	2	2
Total	228		193	185	172	161	147	138	128

+ Less than 0.5 Gg

Note: Totals may not sum due to independent rounding.

* "Other" includes snowmobiles and other recreational equipment, logging equipment, lawn and garden equipment, railroad equipment, airport equipment, commercial equipment, and industrial equipment.

Table 3-25: N₂O Emissions from Mobile Combustion (Gg)

Fuel Type/Vehicle Type	1990		1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	130		166	167	164	158	145	134	123
Passenger Cars	82		86	86	84	80	74	70	64
Light-Duty Trucks	46		76	76	76	74	66	60	54
Heavy-Duty Vehicles	2		4	4	4	4	4	5	5
Motorcycles	+		+	+	+	+	+	+	+
Diesel Highway	1		1	1	1	1	1	1	1
Passenger Cars	+		+	+	+	+	+	+	+

Light-Duty Trucks	+	+	+	+	+	+	+	+
Heavy-Duty Vehicles	1	1	1	1	1	1	1	1
Alternative Fuel Highway	+	+	+	+	+	+	+	+
Non-Highway	10	11	11	11	12	11	12	12
Ships and Boats	1	1	1	1	2	1	2	1
Locomotives	1	1	1	1	1	1	1	1
Farm Equipment	6	6	6	6	6	6	6	5
Construction Equipment	1	1	1	1	1	1	1	1
Aircraft	1	1	1	1	1	1	1	1
Other*	1	1	1	1	1	1	1	1
Total	141	178	179	176	171	158	147	136

+ Less than 0.5 Gg

Note: Totals may not sum due to independent rounding.

* "Other" includes snowmobiles and other recreational equipment, logging equipment, lawn and garden equipment, railroad equipment, airport equipment, commercial equipment, and industrial equipment.

Mobile sources comprise the single largest source category of CO, NO_x, and NMVOC emissions in the United States. In 2003, mobile combustion contributed 89 percent of CO emissions, 56 percent of NO_x emissions, and 46 percent of NMVOC emissions. Since 1990, emissions of NMVOCs from mobile combustion decreased by 42 percent, CO emissions decreased 37 percent, and emissions of NO_x decreased by 14 percent.

Table 3-26: NO_x Emissions from Mobile Combustion (Gg)

Fuel Type/Vehicle Type	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	5,746	4,268	4,090	3,924	3,812	3,715	3,519	3,527
Passenger Cars	3,847	2,447	2,316	2,158	2,084	2,027	1,920	1,924
Light-Duty Trucks	1,364	1,334	1,294	1,268	1,303	1,285	1,217	1,220
Heavy-Duty Vehicles	515	475	467	485	411	390	369	370
Motorcycles	20	13	13	13	13	14	13	13
Diesel Highway	2,956	3,708	3,729	3,671	3,803	3,338	3,162	3,169
Passenger Cars	39	13	11	10	7	6	6	6
Light-Duty Trucks	20	10	9	8	6	5	5	5
Heavy-Duty Vehicles	2,897	3,685	3,709	3,653	3,791	3,326	3,151	3,158
Alternative Fuel Highway^a	IE	IE	IE	IE	IE	IE	IE	IE
Non-Highway	3,432	3,792	3,772	3,705	3,780	3,770	3,707	3,722
Ships and Boats	953	963	919	818	966	971	954	958
Locomotives	857	962	973	910	908	907	891	895
Farm Equipment	63	75	83	84	80	73	72	72
Construction Equipment	437	487	487	497	484	480	472	474
Aircraft ^b	641	708	706	765	697	690	678	681
Other ^c	480	597	604	632	645	650	639	642
Total	12,134	11,768	11,592	11,300	11,395	10,823	10,389	10,418

IE (Included Elsewhere)

Note: Totals may not sum due to independent rounding.

^a NO_x emissions from alternative fuel highway vehicles are included under gasoline and diesel highway vehicles.

^b Aircraft estimates include only emissions related to landing and take-off (LTO) cycles, and therefore do not include cruise altitude emissions.

^c "Other" includes gasoline- and diesel-powered recreational, industrial, lawn and garden, light commercial, logging, airport service, and other equipment.

Table 3-27: CO Emissions from Mobile Combustion (Gg)

Fuel Type/Vehicle Type	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	98,328	67,509	65,246	61,210	60,657	56,716	55,541	52,544
Passenger Cars	60,757	36,825	35,686	32,921	32,867	31,600	30,945	29,275
Light-Duty Trucks	29,237	25,748	24,754	23,343	24,532	22,574	22,107	20,914
Heavy-Duty Vehicles	8,093	4,787	4,642	4,782	3,104	2,411	2,361	2,234

Motorcycles	240	150	163	164	154	131	129	122
Diesel Highway	1,696	1,301	1,202	1,122	1,088	869	851	805
Passenger Cars	35	13	10	10	7	6	6	5
Light-Duty Trucks	22	13	12	9	6	5	5	5
Heavy-Duty Vehicles	1,639	1,276	1,179	1,103	1,075	858	840	795
Alternative Fuel Highway^a	IE	IE	IE	IE	IE	IE	IE	IE
Non-Highway	19,459	21,474	21,493	21,152	21,935	22,387	22,181	22,177
Ships and Boats	1,679	1,948	1,943	2,121	1,946	1,952	1,934	1,934
Locomotives	85	89	83	98	90	90	89	89
Farm Equipment	217	250	274	285	245	233	231	231
Construction Equipment	582	636	633	630	626	621	615	615
Aircraft ^b	1,090	1,098	1,081	1,074	1,047	1,041	1,032	1,031
Other ^c	15,807	17,453	17,478	16,943	17,981	18,449	18,280	18,276
Total	119,482	90,284	87,940	83,484	83,680	79,972	78,574	75,526

IE = Included Elsewhere

Note: Totals may not sum due to independent rounding.

^a CO emissions from alternative fuel highway vehicles are included under gasoline and diesel highway vehicles.

^b Aircraft estimates include only emissions related to landing and take-off (LTO) cycles, and therefore do not include cruise altitude emissions.

^c "Other" includes gasoline- and diesel-powered recreational, industrial, lawn and garden, light commercial, logging, airport service, and other equipment.

Table 3-28: NMVOC Emissions from Mobile Combustion (Gg)

Fuel Type/Vehicle Type	1990	1997	1998	1999	2000	2001	2002	2003
Gasoline Highway	8,110	5,167	5,067	4,924	4,615	4,285	3,931	3,832
Passenger Cars	5,120	2,928	2,895	2,810	2,610	2,393	2,195	2,140
Light-Duty Trucks	2,374	1,882	1,812	1,734	1,750	1,664	1,527	1,488
Heavy-Duty Vehicles	575	336	335	351	232	206	189	184
Motorcycles	42	22	25	28	23	22	20	20
Diesel Highway	406	263	249	230	216	207	190	185
Passenger Cars	16	6	5	5	3	3	3	3
Light-Duty Trucks	14	8	7	6	4	4	3	3
Heavy-Duty Vehicles	377	249	237	219	209	201	184	179
Alternative Fuel Highway^a	IE	IE	IE	IE	IE	IE	IE	IE
Non-Highway	2,416	2,498	2,427	2,432	2,398	2,379	2,438	2,333
Ships and Boats	608	766	763	769	744	730	748	716
Locomotives	33	35	33	38	35	35	36	35
Farm Equipment	28	32	35	38	24	19	20	19
Construction Equipment	85	83	81	81	76	72	74	71
Aircraft ^b	149	142	137	141	130	125	128	123
Other ^c	1,513	1,441	1,378	1,366	1,390	1,397	1,432	1,370
Total	10,933	7,928	7,742	7,586	7,230	6,872	6,560	6,351

IE (Included Elsewhere)

Note: Totals may not sum due to independent rounding.

^a NMVOC emissions from alternative fuel highway vehicles are included under gasoline and diesel highway vehicles.

^b Aircraft estimates include only emissions related to landing and take-off (LTO) cycles, and therefore do not include cruise altitude emissions.

^c "Other" includes gasoline- and diesel-powered recreational, industrial, lawn and garden, light commercial, logging, airport service, and other equipment.

Methodology

Estimates of CH₄ and N₂O emissions from mobile combustion were calculated by multiplying emission factors by measures of activity for each fuel and vehicle type (e.g., light-duty gasoline trucks). Depending upon the category, activity data included such information as fuel consumption, and vehicle miles traveled (VMT). The activity data

and emission factors used are described in the subsections that follow. A complete discussion of the methodology used to estimate emissions from mobile combustion and the emission factors used in the calculations is provided in Annex 3.2.

EPA (2003) provided emissions estimates of NO_x, CO, and NMVOCs for eight categories of highway vehicles,⁴⁴ aircraft, and seven categories of non-highway vehicles.⁴⁵ These emission estimates were provided from preliminary EPA data, which, in its final iteration, will be published on the National Emission Inventory (NEI) Air Pollutant Emission Trends web site. The methodology used to develop these estimates can be found on EPA's Air Pollutant Emission Trends website, at <<http://www.epa.gov/ttn/chief/trends/index.html>>.

Highway Vehicles

Emission estimates for gasoline and diesel highway vehicles were based on VMT and emission factors by vehicle type, fuel type, model year, and control technology. Emissions from alternative fuel vehicles (AFVs)⁴⁶ were based on VMT by vehicle and fuel type.

Emission factors for gasoline and diesel highway vehicles were developed by ICF (2004). These factors were based on EPA and California Air Resources Board (CARB) laboratory test results of different vehicle and control technology types. The EPA and CARB tests were designed following the Federal Test Procedure (FTP), which covers three separate driving segments, since vehicles emit varying amounts of GHGs depending on the driving segment. These driving segments are: (1) a transient driving cycle that includes cold start and running emissions, (2) a cycle that represents running emissions only, and (3) a transient driving cycle that includes hot start and running emissions. For each test run, a bag was affixed to the tailpipe of the vehicle and the exhaust was collected; the content of this bag was then analyzed to determine quantities of gases present. The emission characteristics of segment 2 was used to define running emissions, and subtracted from the total FTP emissions to determine start emissions. These were then recombined based upon the ratio of start to running emissions for each vehicle class from MOBILE6.2 to approximate average driving characteristics.

Emission factors for AFVs were developed after consulting a number of sources, including Argonne National Laboratory's GREET 1.5—Transportation Fuel Cycle Model (Wang 1999), Lipman and Delucchi (2002), the Auto/Oil Air Quality Improvement Research Program (CRC 1997), the California Air Resources Board (Brasil and McMahon 1999), and the University of California Riverside (Norbeck, et al., 1998). The primary approach taken was to calculate CH₄ emissions from actual test data and determine N₂O emissions from NO_x emissions from the same tests. While the formation of N₂O is highly dependent on the type of catalyst used and the catalyst temperature, tailpipe N₂O is likely to increase as engine out NO_x emissions increase. Thus, as a first approximation, the NO_x to N₂O emission ratio will likely be constant for a given emission control group. A complete discussion of the data source and methodology used to determine emission factors from AFVs is provided in Annex 3.2.

Annual VMT data for 1990 through 2003 were obtained from the Federal Highway Administration's (FHWA) Highway Performance Monitoring System database as reported in *Highway Statistics* (FHWA 1996 through 2004). VMT was then allocated from FHWA's vehicle categories to fuel-specific vehicle categories based on estimates of fuel consumption by fuel type for each vehicle category, developed using information on shares of vehicle fuel use for each vehicle category by fuel type reported in DOE (2004) and information on total motor vehicle fuel

⁴⁴ These categories included: gasoline passenger cars, diesel passenger cars, light-duty gasoline trucks less than 6,000 pounds in weight, light-duty gasoline trucks between 6,000 and 8,500 pounds in weight, light-duty diesel trucks, heavy-duty gasoline trucks and buses, heavy-duty diesel trucks and buses, and motorcycles.

⁴⁵ These categories included: locomotives, marine vessels, farm equipment, construction equipment, other off-highway liquid fuel (e.g. recreational vehicles and lawn and garden equipment), and other off-highway gaseous fuel (e.g., other off-highway equipment running on compressed natural gas).

⁴⁶ Alternative fuel and advanced technology vehicles are those that can operate using a motor fuel other than gasoline or diesel. This includes electric or other bifuel or dual fuel vehicles that may be partially powered by gasoline or diesel.

consumption by fuel type from FHWA (1996 to 2004). VMT for AFVs were taken from Browning (2003). The age distributions of the U.S. vehicle fleet were obtained from EPA (2004b) and EPA (2000), and the average annual age-specific vehicle mileage accumulation of U.S. vehicles were obtained from EPA (2000).

Control technology and standards data for highway vehicles were obtained from the EPA's Office of Transportation and Air Quality (EPA 2004a, 2004d, 2000, 1998, and 1997). These technologies and standards are defined in Annex 3.2, and were compiled from EPA (1993), EPA (1994a), EPA (1994b), EPA (1998), EPA (1999), and IPCC/UNEP/OECD/IEA (1997).

Preliminary estimates for NO_x, CO, and NMVOCs were obtained from EPA (2004e) and disaggregated based on EPA (2003), which, in its final iteration, will be published on the National Emission Inventory (NEI) Air Pollutant Emission Trends web site.

Non-Highway

Fuel consumption data were employed as a measure of activity for non-highway vehicles, and then fuel-specific emission factors were applied.⁴⁷ Activity data were obtained from AAR (2004), BEA (1991 through 2004), Benson (2002 through 2004), DOE (1993 through 2004), DESC (2004), DOC (1991 through 2004), DOT (1991 through 2004), EIA (2002a), EIA (2002b), EIA (2004a), EIA (2004b), EIA (2003 through 2004), EIA (1991 through 2004), EPA (2004c), and FAA (2004). Emission factors for non-highway modes were taken from IPCC/UNEP/OECD/IEA (1997).

Uncertainty

This section discusses the uncertainty of the emissions estimates for CH₄ and N₂O. Uncertainty was analyzed separately for highway vehicles and non-highway vehicles, due to differences in their characteristics and their contributions to total mobile source emissions.

Uncertainty analyses were not conducted for CO, NO_x, or NMVOC emissions. Emission factors for these gases have been extensively researched, since these gases are regulated emissions from motor vehicles in the United States, and the uncertainty of these emissions estimates is believed to be relatively low. A much higher level of uncertainty is associated with CH₄ and N₂O emission factors, since emissions of these gases are not regulated in the United States, and unlike CO₂ emissions, the emission pathways of CH₄ and N₂O are also highly complex.

Highway Vehicles

A quantitative uncertainty analysis was conducted for the highway portion of the mobile source sector using the IPCC-recommended Tier 2 uncertainty estimation methodology, Monte Carlo Simulation technique, using @RISK software. The uncertainty analysis was performed on 2003 estimates of CH₄ and N₂O emissions, incorporating probability distribution functions associated with the major input variables. For the purposes of this analysis, the uncertainty was modeled for the following two major sets of input variables: (1) vehicle mile traveled (VMT) data, by vehicle and fuel type and (2) emission factor data, by vehicle, fuel, and control technology type.

Mobile combustion emissions of CH₄ and N₂O per vehicle mile traveled vary significantly due to fuel type and composition, technology type, operating speeds and conditions, type of emission control equipment, equipment age, and operating and maintenance practices. The primary activity data, VMT, are collected and analyzed each year by government agencies.

⁴⁷ The consumption of international bunker fuels is not included in these activity data, but is estimated separately under the International Bunker Fuels source category.

To determine the uncertainty associated with the activity data used in the calculations of CH₄ and N₂O emissions, the agencies and the experts that supply the data were contacted. Because few of these sources were able to provide quantitative estimates of uncertainty, expert quantitative judgments were used to assess the uncertainty associated with the activity data.

The emission factors for highway vehicles used in the Inventory were obtained from ICF (2004). These factors were based on laboratory testing of vehicles. While the controlled testing environment simulates real driving conditions, emission results from such testing can only approximate real world conditions and emissions. For some vehicle and control technology types, the testing did not yield statistically significant results within the 95 percent confidence interval, requiring expert judgments to be used in developing the emission factors. In those cases, the emission factors were developed based on comparisons of fuel consumption between similar vehicle and control technology categories.

The estimates of VMT for highway vehicles by vehicle type in the United States were provided by FHWA (1996 through 2004), and were generated through the cooperation of FHWA and state and local governments. These estimates are subject to several possible sources of error, such as unregistered vehicles, unreported fuel sales to avoid fuel taxes, differences in achieved versus estimated fuel economy, and measurement and estimation errors. These VMT were apportioned by fuel type, and then allocated to individual model years using temporal profiles of both the vehicle fleet by age and vehicle usage by model year in the United States provided by EPA (2004b) and EPA (2000). While the uncertainty associated with total U.S. VMT is believed to be low, the uncertainty within individual source categories was assumed to be higher given uncertainties associated with apportioning total VMT into individual vehicle categories, by technology type, and equipment age. The uncertainty of individual estimates was assumed to relate to the magnitude of estimated VMT (i.e., it was assumed smaller sources had greater percentage uncertainty). A further source of uncertainty occurs since FHWA and EPA use different definitions of vehicle type and estimates of VMT by vehicle type (provided by FHWA) are broken down by fuel type using EPA vehicle categories.

A total of 69 highway data input variables were simulated through Monte Carlo Simulation technique using @RISK software. Variables included VMT and emission factors for individual vehicle categories and technologies. In developing the uncertainty estimation model, a normal distribution was assumed for all activity-related input variables (e.g., VMT) except in the case of buses, in which a triangular distribution was used. The dependencies and other correlations among the activity data were incorporated into the model to ensure consistency in the model specification and simulation. Emission factors were assigned uniform distributions, with upper and lower bounds assigned to input variables based on 97.5 percent confidence intervals of laboratory test data. In cases where data did not yield statistically significant results within the 95 percent confidence interval, estimates of upper and lower bounds were made using expert judgment. The bounds for the emission factor-related input variables were typically asymmetrical around their inventory estimates. Bias (or systematic uncertainties) associated with the emission factors was incorporated into the analysis when expert judgments were applied to the laboratory test results in determining the uncertainty characteristics and/or the bounds of the emission factors.⁴⁸ The results of this analysis are reported in the section below, titled *Quantitative Estimates of Uncertainty*.

Non-Highway

Emissions from non-highway vehicles are a small portion of total emissions from mobile sources, representing 20 percent of CH₄ emissions from mobile sources and 9 percent of N₂O emissions from mobile sources in 2003. Since they comprise a small share of mobile source emissions, even large uncertainties in these estimates would have a relatively small impact on the total emission estimate for mobile sources. As a result, a quantitative analysis of uncertainty of emissions from non-highway vehicles has not been performed. However, sources of uncertainty for

⁴⁸ Random uncertainties are the main focus of statistical uncertainty analysis. Uncertainty estimates elicited from experts include both random and systematic uncertainty. Hence, both these types of uncertainty are represented in this uncertainty analysis.

non-highway vehicles are being investigated by examining the underlying uncertainty of emission factors and fuel consumption data.

Overall, a significant amount of uncertainty is associated with the emission estimates for non-road sources. A primary cause is a large degree of uncertainty surrounding emission factors. The *IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories* reports that CH₄ emissions from aviation and marine sources may be uncertain by a factor of two, while N₂O emissions may be uncertain by an order of magnitude for marine sources and several orders of magnitude for aviation. No information is provided on the uncertainty of emission factors for other non-highway sources.

Fuel consumption data have a lower uncertainty than emission factors, though large uncertainties do exist for individual sources. Fuel consumption for off-highway vehicles (i.e., equipment used for agriculture, construction, lawn and garden, railroad, airport ground support, etc., as well as recreational vehicles) was generated by EPA's NONROAD model (EPA 2004c). This model estimates fuel consumption based on estimated equipment/vehicle use (in hours) and average fuel consumed per hour of use. Since the fuel estimates are not based upon documented fuel sales or consumption, a fair degree of uncertainty accompanies these estimates.

Distillate consumption for ships and boats was obtained from sales estimates from *EIA's Fuel Oil and Kerosene Sales* (EIA 1991 through 2004). The estimates for distillate consumption have associated uncertainty, as EIA's estimates are based on sales to economic sectors, and it can be difficult to determine how much of the fuel sold in each sector is used by mobile or stationary sources and to further attribute this consumption to specific final users. For example, some fuel purchased by the marine sector may be used for operating heavy equipment or even generators, instead of being used entirely by ships and boats. This distinction between mobile and stationary fuel users is not made by EIA.

EIA does provide coefficients of variation to estimate sampling error, which occurs due to the fact that these estimates are based on a sample set. However, as EIA points out, these coefficients do not take into account all the sources of potential bias, which includes incomplete information, misinterpretation of survey questions, and other factors that may cause estimates of fuel sales to be different from actual sales. In addition, diesel for ships and boats is adjusted for bunker fuel consumption, which introduces an additional (and much higher) level of uncertainty.

Domestic consumption for residual fuel consumption by ships and boats is obtained from EIA (2004a). These estimates fluctuate widely from year to year. Such fluctuations cannot be fully explained without further analysis of the underlying activity data sets and, as such, the estimates are believed to be highly uncertain. The estimate of domestic consumption is then adjusted downward to account for international bunker fuels, which represents the primary use of residual fuel by ships and boats. As the international bunker fuel data are considered to have a moderate level of uncertainty,⁴⁹ the overall uncertainty of the domestic ships and boats estimate for residual fuel consumption is considered high.

Domestic jet fuel and aviation gasoline consumption data are obtained from EIA (2004a). Like diesel and residual marine fuel consumption, jet fuel consumption for aviation is adjusted downward to account for international bunker fuels. The international bunker fuel estimates introduce a significant amount of uncertainty. Additionally, all jet fuel consumption in the transportation sector is assumed to be consumed by aircraft. Some fuel purchased by airlines is not used in aircraft, but instead used to power auxiliary power units, in ground equipment, and to test engines. Some jet fuel may also be used for other purposes such as blending with diesel fuel or heating oil.

In calculating CH₄ emissions from aircraft, an average emission factor is applied to total jet fuel consumption. This average emission factor takes into account the fact that CH₄ emissions occur only during the landing and take-off

⁴⁹ This is discussed in the section on International Bunker Fuels.

(LTO) cycles, with no CH₄ being emitted during the cruise cycle. However, a better approach would be to apply emission factors based on the number of LTO cycles.

Finally, U.S. aircraft emission estimates for CO, NO_x, and NMVOCs from EPA (2004e and 2003) are based on LTO cycles and only estimate near ground-level emissions, which are more relevant for air quality evaluations. These estimates also include both domestic and international flights. Therefore, estimates presented here may overestimate IPCC-defined domestic CO, NO_x, and NMVOC emissions by including LTO cycles by aircraft on international flights but underestimate total emissions because they exclude emissions from aircraft on domestic flight segments at cruising altitudes.

Quantitative Estimates of Uncertainty

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-29. Mobile combustion CH₄ emissions in 2003 were estimated to be between 2.5 and 2.8 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 9 percent below to 4 percent above the 2003 emission estimate of 2.7 Tg CO₂ Eq. Also at a 95 percent confidence level, mobile combustion N₂O emissions in 2003 were estimated to be between 35.2 and 52.8 Tg CO₂ Eq., indicating a range of 16 percent below to 26 percent above the 2003 emission estimate of 42.1 Tg CO₂ Eq.

Table 3-29: Tier 2 Quantitative Uncertainty Estimates for CH₄ and N₂O Emissions from Mobile Sources (Tg CO₂ Eq. and Percent)

Source	Gas	2003 Emission Estimate (Tg CO ₂ Eq.)	Uncertainty Range Relative to Emission Estimate ^a			
			(Tg CO ₂ Eq.)		(%)	
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Mobile Sources	CH ₄	2.7	2.5	2.8	-9%	+4%
Mobile Sources	N ₂ O	42.1	35.2	52.8	-16%	+26%

^aRange of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

This uncertainty analysis is a continuation of a multi-year process for developing credible quantitative uncertainty estimates for this source category using the IPCC Tier 2 approach to uncertainty analysis. In the upcoming years, the type and the characteristics of the actual probability density functions underlying the input variables will be identified and more credibly characterized. Accordingly, the quantitative uncertainty estimates reported in this section should be considered as preliminary and illustrative.

QA/QC and Verification

A source-specific QA/QC plan for mobile combustion was developed and implemented. This effort included a Tier 1 analysis, as well as portions of a Tier 2 analysis. The Tier 2 procedures focused on the emission factor and activity data sources, as well as the methodology used for estimating emissions. Because significant changes were made to highway vehicle emission factors and non-highway fuel consumption estimates (see Recalculations Discussion below), QA/QC efforts were focused in particular on emissions impacted by these changes. These procedures included a qualitative assessment of the emission estimates to determine whether they appear consistent with the most recent activity data and emission factors available. A comparison of historical emissions between this year's and last year's Inventories was also conducted, and was qualitatively assessed to ensure that the changes in estimates were consistent with the changes in activity data and emission factors.

Recalculations Discussion

In order to ensure the highest quality estimates, the methodology is continuously revised based on comments from internal and external reviewers. This year, adjustments were made to emission factors and activity data to incorporate new research and additional data sources.

The most significant changes in this year's Inventory are the revisions to the highway emission factors for CH₄ and N₂O. The previous emission factors had been derived using methodology similar to that outlined in IPCC/UNEP/OECD/IEA (1997). However, preliminary tests suggested that these emission factors may not have been representative of actual emissions. EPA sponsored laboratory testing that resulted in the new emission factors used in this report. The emission factors changed notably, ranging from a decrease of 97 percent to an increase of 114 percent for individual vehicle/fuel type/technology types; the combined effect of these changes in emission factors significantly lowered CH₄ and N₂O emissions from highway vehicles. Given the large contribution of highway vehicles to the mobile totals, these new emission factors were major drivers behind the changes in CH₄ and N₂O estimates. However, because these emission factors do not impact CO₂ emissions, their impact is less significant when considering total mobile source emissions of all GHGs.

Additionally, the methodology for estimating VMT by vehicle/fuel type category (e.g., LDGV, LDDV, LDGT, LDDT, etc.) was revised. The purpose of this revision was to ensure that the VMT estimates for individual vehicle categories match up with VMT estimates by vehicle type from FHWA (1996 through 2004), which is the recognized source of national VMT estimates. The new methodology apportions FHWA's estimate of VMT for each vehicle category (e.g., light-duty vehicles, light-duty trucks, buses, heavy-duty trucks) to fuel-based vehicle categories based on estimates of gasoline and diesel fuel consumption for each vehicle category, drawn from DOE (2004). This change had a small impact on emissions.

The vehicle age distribution values for highway vehicles were also revised. Previously, the Inventory relied on one vehicle age distribution across 25 model years from EPA (2000), which was applied for each year. For the current Inventory, that same distribution was applied for years 1990 through 1998, and annually variable vehicle age distributions were applied for years 1999 through 2003 based on data obtained from EPA's MOVES model (EPA 2004b). The annually variable age distributions more accurately reflect changes in the vehicle stock, and include 31 model years.

Emissions from gasoline- and diesel-electric hybrid vehicles are now included under gasoline and diesel vehicles instead of alternative fuel/advanced technology category, since these vehicles only run on traditional motor gasoline. This change did not have an impact on total emissions.

EPA's NONROAD model is now used as the primary data source for fuel consumption by off-highway equipment (e.g., construction, agricultural, lawn and garden equipment), rather than data from FHWA's *Highway Statistics* (FHWA 1996 through 2004), EIA's *Fuel Oil and Kerosene Sales* (EIA 1991 through 2004), and several other sources. The NONROAD model is also used to estimate fuel consumption by recreational boats, rather than data from FHWA's *Highway Statistics*. Some of the individual sources used in the previous Inventory had not been updated for several years, while others had updated their methodologies, resulting in large variations in estimates from year to year. Use of the NONROAD Model for all of these estimates is believed to provide more accurate, up-to-date, and consistent estimates.

Other changes include minor revisions to historical aircraft fuel consumption estimates in FAA (2004) and rail consumption provided from Benson (2004). These revisions were not significant, and the overall impact on emissions estimates is small.

Overall, these changes resulted in an average annual decrease of 0.6 Tg CO₂ Eq. (14 percent) in CH₄ emissions from mobile combustion and an average annual decrease of 6.2 Tg CO₂ Eq. (11 percent) in N₂O emissions from mobile combustion for the period 1990 through 2002.

Planned Improvements

While the data used for this report represent the most accurate information available, three areas have been identified that could potentially be improved in the short term given resource availability:

- 1) *Reconcile Fuel Consumption Estimates used for Calculating N₂O/CH₄ and CO₂* – Estimates of transportation fuel consumption by fuel type from EIA are used as the basis for estimating CO₂ emissions from the transportation sector. These estimates are then apportioned to mode and vehicle category based on "bottom up" estimates of fuel

consumption from sources such as FHWA's *Highway Statistics* (FHWA 1996 through 2004) and DOE's *Transportation Energy Data Book* (DOE 1993 through 2004). These sources are also used to develop N₂O and CH₄ estimates. The EIA fuel consumption estimates, however, differ from the estimates derived using "bottom up" sources. For certain vehicle categories this leads to CO₂ emissions trends that conflict with those of the "bottom up" sources. Potential improvements include reconciling fuel consumption estimates from EIA and other data sources, and revising the current process of allocating CO₂ emissions to reflect trends from the more detailed vehicle category estimates of fuel consumption.

2) *Improve consideration of emissions from trucks used off-road* – Some light- and heavy-duty trucks travel for a portion of their mileage off-road. N₂O and CH₄ estimates for highway vehicles are developed based on vehicle mileage data from FHWA's *Highway Statistics*, which in turn, are drawn from the Highway Performance Monitoring System (HPMS). These emissions estimates do not address travel by trucks off-road. Gasoline fuel consumed by trucks used off-road for construction, agriculture, and other industrial/commercial uses is reported in *Highway Statistics*, and is included as part of the non-road agriculture and construction categories. However, diesel fuel consumed by trucks used off-road is not addressed in the current Inventory, and further work should be conducted to develop estimates of off-road truck use of diesel fuel. In addition, default emission factors from IPCC are applied to the off-highway modes. As a result, the emissions factors for agricultural equipment are applied both to equipment and trucks used in agriculture, and emissions factors for construction equipment are applied both to equipment and trucks used in construction. Emission estimates would be improved through an investigation of more appropriate emission factors for off-road trucks.

3) *Improve estimation of VMT by vehicle/fuel type category* – The current Inventory process for estimating VMT by vehicle/fuel type category involves apportioning VMT by vehicle type to each fuel type on the basis of fuel consumption. While this is a reasonable simplification, this approach implicitly assumes the same average fuel economy for gasoline and diesel vehicles. A more accurate apportionment for VMT by fuel type for light-duty trucks and medium/heavy-duty trucks could potentially be developed using data on vehicle travel from the Vehicle Inventory and Use Survey and other publications, or using VMT breakdowns by vehicle/fuel type combinations from the MOBILE6 or MOVES models. These sources should be investigated in order to develop a more robust apportionment method.

3.5. Coal Mining (IPCC Source Category 1B1a)

Three types of coal mining related activities release CH₄ to the atmosphere: underground mining, surface mining, and post-mining (i.e., coal-handling) activities. Underground coal mines contribute the largest share of CH₄ emissions. All 101 gassy underground coal mines employ ventilation systems to ensure that CH₄ levels remain within safe concentrations. These systems can exhaust significant amounts of CH₄ to the atmosphere in low concentrations. Additionally, twenty-one U.S. coal mines supplement ventilation systems with degasification systems. Degasification systems are wells drilled from the surface or boreholes drilled inside the mine that remove large volumes of CH₄ before, during, or after mining. In 2003, ten coal mines collected CH₄ from degasification systems and sold this gas to a pipeline, thus reducing emissions to the atmosphere. In addition, one coal mine used CH₄ from its degasification system to heat mine ventilation air on site. Surface coal mines also release CH₄ as the overburden is removed and the coal is exposed, but the level of emissions is much lower than from underground mines. Finally, some of the CH₄ retained in the coal after mining is released during processing, storage, and transport of the coal.

Total CH₄ emissions in 2003 were estimated to be 53.8 Tg CO₂ Eq. (2,561 Gg), a decline of 34 percent since 1990 (see Table 3-30 and Table 3-31). Of this amount, underground mines accounted for 70 percent, surface mines accounted for 16 percent, and post-mining emissions accounted for 14 percent. In 1993, CH₄ generated from underground mining dropped, primarily due to labor strikes at many large underground mines. In 1994 and 1995, CH₄ emissions increased due to resumed production at high emitting mines after the labor strike. The decline in CH₄ emissions from underground mines from 1996 to 2002 was the result of the reduction of overall coal production, the mining of less gassy coal, and an increase in CH₄ recovered and used. CH₄ emissions increased slightly in 2003 due to additional gas drainage being vented to the atmosphere and a reduction in CH₄ recovery. Surface mine emissions and post-mining emissions remained relatively constant from 1990 to 2003.

Table 3-30: CH₄ Emissions from Coal Mining (Tg CO₂ Eq.)

Activity	1990	1996	1997	1998	1999	2000	2001	2002	2003
Underground Mining	62.1	45.3	44.3	44.4	41.6	39.4	38.1	35.8	37.6
Liberated	67.6	59.8	55.7	58.6	54.4	54.0	54.2	53.3	53.6
Recovered & Used	(5.6)	(14.5)	(11.4)	(14.2)	(12.7)	(14.6)	(16.1)	(17.5)	(16.1)
Surface Mining	10.4	9.2	9.3	9.4	9.0	8.8	9.2	8.8	8.4
Post-Mining (Underground)	7.7	7.2	7.4	7.4	6.8	6.7	6.8	6.4	6.4
Post-Mining (Surface)	1.7	1.5	1.5	1.5	1.5	1.4	1.5	1.4	1.4
Total	81.9	63.2	62.6	62.8	58.9	56.2	55.6	52.4	53.8

Note: Totals may not sum due to independent rounding.

Table 3-31: CH₄ Emissions from Coal Mining (Gg)

Activity	1990	1996	1997	1998	1999	2000	2001	2002	2003
Underground Mining	2,956	2,158	2,111	2,117	1,982	1,876	1,816	1,705	1,788
Liberated	3,220	2,850	2,654	2,791	2,589	2,573	2,580	2,538	2,554
Recovered & Used	(265)	(692)	(543)	(674)	(607)	(697)	(765)	(833)	(766)
Surface Mining	497	438	445	448	428	417	438	420	402
Post-Mining (Underground)	367	341	354	352	325	317	323	304	305
Post-Mining (Surface)	81	71	72	73	69	68	71	68	65
Total	3,900	3,008	2,983	2,989	2,805	2,677	2,647	2,497	2,561

Note: Totals may not sum due to independent rounding.

Methodology

The methodology for estimating CH₄ emissions from coal mining consists of two parts. The first part involves estimating CH₄ emissions from underground mines. Because of the availability of ventilation system measurements, underground mine emissions can be estimated on a mine-by-mine basis and then summed to determine total emissions. The second step involves estimating emissions from surface mines and post-mining activities by multiplying basin-specific coal production by basin-specific emission factors.

Underground mines. Total CH₄ emitted from underground mines was estimated as the sum of CH₄ liberated from ventilation systems and CH₄ liberated by means of degasification systems, minus CH₄ recovered and used. The Mine Safety and Health Administration (MSHA) samples CH₄ emissions from ventilation systems for all mines with detectable⁵⁰ CH₄ concentrations. These mine-by-mine measurements are used to estimate CH₄ emissions from ventilation systems.

Some of the higher-emitting underground mines also use degasification systems (e.g., wells or boreholes) that remove CH₄ before, during, or after mining. This CH₄ can then be collected for use or vented to the atmosphere. Various approaches were employed to estimate the quantity of CH₄ collected by each of the twenty-one mines using these systems, depending on available data. For example, some mines report to EPA the amount of CH₄ liberated from their degasification systems. For mines that sell recovered CH₄ to a pipeline, pipeline sales data published by state petroleum and natural gas agencies were used to estimate degasification emissions. For those mines for which no other data are available, default recovery efficiency values were developed, depending on the type of degasification system employed.

Finally, the amount of CH₄ recovered by degasification systems and then used (i.e., not vented) was estimated. This calculation was complicated by the fact that most CH₄ is not recovered and used during the same year in which the particular coal seam is mined. In 2003, ten active coal mines sold recovered CH₄ into the local gas pipeline

⁵⁰ MSHA records coal mine methane readings with concentrations of greater than 50 ppm (parts per million) methane. Readings below this threshold are considered non-detectable.

networks, while one coal mine used recovered CH₄ on site. Emissions avoided for these projects were estimated using gas sales data reported by various state agencies. For most mines with recovery systems, companies and state agencies provided individual well production information, which was used to assign gas sales to a particular year. For the few remaining mines, coal mine operators supplied information regarding the number of years in advance of mining that gas recovery occurs.

Surface Mines and Post-Mining Emissions. Surface mining and post-mining CH₄ emissions were estimated by multiplying basin-specific coal production, obtained from the Energy Information Administration's *Coal Industry Annual* (see Table 3-32) (EIA 2003), by basin-specific emission factors. Surface mining emission factors were developed by assuming that surface mines emit two times as much CH₄ as the average *in situ* CH₄ content of the coal. Revised data on *in situ* CH₄ content and emissions factors are taken from EPA (1996) and AAPG (1984). This calculation accounts for CH₄ released from the strata surrounding the coal seam. For post-mining emissions, the emission factor was assumed to be 32.5 percent of the average *in situ* CH₄ content of coals mined in the basin.

Table 3-32: Coal Production (Thousand Metric Tons)

Year	Underground	Surface	Total
1990	384,250	546,818	931,068
1991	368,635	532,656	901,291
1992	368,627	534,290	902,917
1993	318,478	539,214	857,692
1994	362,065	575,529	937,594
1995	359,477	577,638	937,115
1996	371,816	593,315	965,131
1997	381,620	607,163	988,783
1998	378,964	634,864	1,013,828
1999	355,433	642,877	998,310
2000	338,173	635,592	973,765
2001	345,305	676,142	1,021,446
2002	324,219	667,619	991,838
2003	320,047	651,251	971,297

Uncertainty

The emission estimates from underground ventilation systems were based on actual measurement data, which are believed to have relatively low uncertainty. A degree of imprecision was introduced because the measurements were not continuous but rather an average of quarterly instantaneous readings. Additionally, the measurement equipment used possibly resulted in an average of 10 percent overestimation of annual CH₄ emissions (Mutmansky and Wang 2000). Estimates of CH₄ liberated and recovered by degasification systems are also relatively certain because many coal mine operators provided information on individual well gas sales and mined through dates. Many of the recovery estimates use data on wells within 100 feet of a mined area. A level of uncertainty currently exists concerning the radius of influence of each well. The number of wells counted, and thus the avoided emissions, may increase if the drainage area is found to be larger than currently estimated.

Compared to underground mines, there is considerably more uncertainty associated with surface mining and post-mining emissions because of the difficulty in developing accurate emission factors from field measurements. However, since underground emissions comprise the majority of total coal mining emissions, the uncertainty associated with underground emissions is the primary factor that determines overall uncertainty. The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-33. Coal mining CH₄ emissions in 2003 were estimated to be between 51.9 and 55.7 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 4 percent below to 4 percent above the 2003 emission estimate of 53.8 Tg CO₂ Eq.

Table 3-33: Tier 2 Quantitative Uncertainty Estimates for CH₄ Emissions from Coal Mining (Tg CO₂ Eq. and Percent)

Source	Gas	2003 Emission	Uncertainty Range Relative to Emission Estimate ^a			
		Estimate	(Tg CO ₂ Eq.)		Lower	
		(Tg CO ₂ Eq.)	(Tg CO ₂ Eq.)		Upper	
			Lower	Upper	Lower	Upper
			Bound	Bound	Bound	Bound
Coal Mining	CH ₄	53.8	51.9	55.7	-4%	+4%

^a Range of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

Recalculations Discussion

In-situ gas content is the principal variable used to determine post-mining methane emissions of mined coal. Previously, in-situ values used were based on average CH₄ content values summarized in Exhibit 3-4 of the U.S. EPA publication, EPA/400/9-90/008; *Methane Emissions From Coal Mining, Issues and Opportunities, September 1990*. The original source of information is derived from three primary sources: 1986 USBM Circular 9067, *Results of the Direct Method Determination of the Gas Contents of U.S. Coal Basins*, 1983 U.S. DOE Report (DOE/METC/83-76), *Methane Recovery from Coalbeds: A Potential Energy Source*, and a series of 1986-88 Gas Research Institute Topical Reports called *A Geologic Assessment of Natural Gas from Coal Seams*. No data was available for eight of the coal mining states and therefore default values from other coal basins were assigned to those states.

Since Circular 9067 contained only a portion of the gas content data compiled by USBM, the complete dataset, published in 1996 *Evaluation and Analysis of Gas Content and Coal Properties of Major Coal Bearing Regions of the United States*, EPA/600/R-96-065, is now the basis of new in-situ gas content value. In addition, gas content data from the U.S. DOE Methane Recovery from Coalbed Projects (MRCP), which was the original source of data for the GRI Topical Reports noted above, was utilized. (Condensed versions of the original MRCP reports for 13 U.S. coal basins are compiled in *Coalbed Methane Resources of the United States, AAPG Studies in Geology Series #17*, published in 1984).

The compiled gas content data for each of the coal basins was sorted by depth to determine in-situ values for surface and underground mines, separately. Overburden depths of surface mines were analyzed using *Keystone Coal Industry Manuals* from 1991 through 2003 and found that the maximum depth was 250 feet. Therefore, gas content data from samples taken less than 250 feet deep were assigned to surface mines and the samples collected from deeper depths to underground mines. The combination of these changes resulted in an average annual decrease of 0.2 Tg CO₂ Eq. (0.3 percent) in CH₄ emissions from coal mining for the period 1990 through 2002.

Planned Improvements

To reduce the uncertainty associated with the radius of influence of each well, the appropriate drainage radius will be investigated for future inventories. Since the number of wells counted may increase if the drainage area is found to be larger than currently estimated, additional mines may be included in future estimates of recovery.

3.6. Abandoned Underground Coal Mines (IPCC Source Category 1B1a)

All underground and surface coal mining liberates CH₄ as part of the normal mining operations. The amount of CH₄ liberated depends on the amount that resides in the coal (“*in situ*”) and surrounding strata when mining occurs. The in-situ CH₄ content depends upon the amount of CH₄ created during the coal formation (i.e., coalification) process, and the geologic characteristics of the coal seams. During coalification, more deeply buried deposits tend to generate more CH₄ and retain more of the gas after uplift to minable depths. Deep underground coal seams generally have higher CH₄ contents than shallow coal seams or surface deposits.

Underground coal mines contribute the largest share of CH₄ emissions, with active underground mines the leading source of underground emissions. However, mines also continue to release CH₄ after closure. As mines mature and coal seams are mined through, mines close and are abandoned. Many are sealed and some flood through intrusion of groundwater or surface water into the void. Shafts or portals are generally filled with gravel and capped with a

concrete seal, while vent pipes and boreholes are plugged in a manner similar to oil and gas wells. Some abandoned mines are vented to the atmosphere to prevent the buildup of CH₄ that may find its way to surface structures through overburden fractures. As work stops within the mines, the CH₄ liberation decreases but it does not stop completely. Following an initial decline, abandoned mines can liberate CH₄ at a near-steady rate over an extended period of time, or, if flooded, produce gas for only a few years. The gas can migrate to the surface through the conduits described above, particularly if they have not been sealed adequately. In addition, diffuse emissions can occur when CH₄ migrates to the surface through cracks and fissures in the strata overlying the coal mine. The following factors influence abandoned mine emissions:

- Time since abandonment;
- Gas content and adsorption characteristics of coal;
- Methane flow capacity of the mine;
- Mine flooding;
- Presence of vent holes; and
- Mine seals.

Gross abandoned mine methane emissions ranged from 6.1 to 9.6 Tg CO₂ Eq. from 1990 through 2003, varying as much as 1.3 Tg CO₂ Eq. from year to year. Fluctuations were due mainly to the number of mines closed during a given year as well as the magnitude of the emissions from those mines when active. Abandoned mine emissions peaked in 1996 (8.6 Tg CO₂ Eq.) due to the large number of mine closures from 1994 to 1996 (70 gassy mines closed during the three-year period). In spite of this rapid rise, abandoned mine emissions have been generally on the decline since 1996 (with slight increases in 1999 and 2000). There were fewer than thirteen gassy mine closures during each of the years from 1998 through 2003. By 2003, abandoned mine emissions were reduced to 6.4 Tg CO₂ Eq. (see Table 3-34 and Table 3-35).

Table 3-34: CH₄ Emissions from Abandoned Coal Mines (Tg CO₂ Eq.)

Activity	1990	1997	1998	1999	2000	2001	2002	2003
Abandoned Underground Mines	6.1	9.6	8.8	9.0	9.3	8.5	8.0	7.9
Recovered & Used	0	1.5	1.7	1.6	1.5	1.5	1.6	1.5
Total	6.1	8.1	7.1	7.3	7.7	6.9	6.4	6.4

Note: Totals may not sum due to independent rounding.

Table 3-35: CH₄ Emissions from Abandoned Coal Mines (Gg)

Activity	1990	1997	1998	1999	2000	2001	2002	2003
Abandoned Underground Mines	288	458	421	426	441	403	380	377
Recovered & Used	-	74	80	78	73	73	77	72
Total	288	385	341	349	369	331	303	306

Note: Totals may not sum due to independent rounding.

Methodology

Estimating CH₄ emissions from an abandoned coal mine requires predicting the emissions of a mine from the time of abandonment through the inventory year of interest. The flow of CH₄ from the coal to the mine void is primarily dependent on mine's emissions when active and the extent to which the mine is flooded or sealed. The CH₄ emission rate before abandonment reflects the gas content of the coal, rate of coal mining, and the flow capacity of the mine in much the same way as the initial rate of a water-free conventional gas well reflects the gas content of the producing formation and the flow capacity of the well. Existing data on abandoned mine emissions through time, although sparse, appear to fit the hyperbolic type of decline curve used in forecasting production from natural gas wells.

In order to estimate CH₄ emissions over time for a given mine, it is necessary to apply a decline function, initiated upon abandonment, to that mine. In the analysis, mines were grouped by coal basin with the assumption that they will generally have the same initial pressures, permeability and isotherm. As CH₄ leaves the system, the reservoir pressure, P_r , declines as described by the isotherm. The emission rate declines because the mine pressure (P_w) is essentially constant at atmospheric pressure, for a vented mine, and the PI term is essentially constant at the

pressures of interest (atmospheric to 30 psia). A rate-time equation can be generated that can be used to predict future emissions. This decline through time is hyperbolic in nature and can be empirically expressed as:

$$q = q_i(1 + bD_it)^{(-1/b)}$$

Where:

- q is the gas rate at time t in mcf/d
- q_i is the initial gas rate at time zero (t_0) in million cubic feet per day (mcf/d)
- b is the hyperbolic exponent, dimensionless
- D_i is the initial decline rate, 1/yr
- t is elapsed time from t_0 in years

This equation is applied to mines of various initial emission rates that have similar initial pressures, permeability and adsorption isotherms (EPA 2003).

The decline curves are also affected by both sealing and flooding. Based on field measurement data, it was assumed that most U.S. mines prone to flooding will become completely flooded within 8 years and therefore no longer have any measurable CH₄ emissions. Based on this assumption, an average decline rate for flooding mines was established by fitting a decline curve to emissions from field measurements. An exponential equation was developed from emissions data measured at eight abandoned mines known to be filling with water located in two of the five basins. Using a least squares, curve-fitting algorithm, emissions data were matched to the exponential equation shown below. There was not enough data to establish basin-specific equations as was done with the vented, non-flooding mines (EPA 2003).

$$q = q_i e^{(-Dt)}$$

Where:

- q is the gas flow rate at time t in mcf/d
- q_i is the initial gas flow rate at time zero (t_0) in mcf/d
- D is the decline rate, 1/yr
- t is elapsed time from t_0 in years

Seals have an inhibiting effect on the rate of flow of CH₄ into the atmosphere compared to the rate that would be emitted if the mine had an open vent. The total volume emitted will be the same, but will occur over a longer period. The methodology, therefore, treats the emissions prediction from a sealed mine similar to emissions from a vented mine, but uses a lower initial rate depending on the degree of sealing. The computational fluid dynamics simulator was again used with the conceptual abandoned mine model to predict the decline curve for inhibited flow. The percent sealed is defined as $100 \times (1 - \text{initial emissions from sealed mine} / \text{emission rate at abandonment prior to sealing})$. Significant differences are seen between 50 percent, 80 percent and 95 percent closure. These decline curves were therefore used as the high, middle, and low values for emissions from sealed mines (EPA 2003).

For active coal mines, those mines producing over 100 mcf/d account for 98 percent of all CH₄ emissions. This same relationship is assumed for abandoned mines. It was determined that 434 abandoned mines closing after 1972 produced emissions greater than 100 mcf/d when active. Further, the status of 256 of the 434 mines (or 59 percent) is known to be either 1) vented to the atmosphere, 2) sealed to some degree (either earthen or concrete seals), or 3) flooded (enough to inhibit methane flow to the atmosphere). The remaining 41 percent of the mines were placed in one of the three categories by applying a probability distribution analysis based on the known status of other mines located in the same coal basin (EPA 2003).

Inputs to the decline equation require the average emission rate and the date of abandonment. Generally this data is available for mines abandoned after 1972; however, such data are largely unknown for mines closed before 1972. Information that is readily available such as coal production by state and county are helpful, but do not provide enough data to directly employ the methodology used to calculate emissions from mines abandoned after 1971. It is

assumed that pre-1972 mines are governed by the same physical, geologic, and hydrologic constraints that apply to post-1972 mines; thus, their emissions may be characterized by the same decline curves.

During the 1970s, 78 percent of CH₄ emissions from coal mining came from seventeen counties in seven states. In addition, mine closure dates were obtained for two states, Colorado and Illinois, throughout the 20th century. The data was used to establish a frequency of mine closure histogram (by decade) and applied to the other five states with gassy mine closures. As a result, basin-specific decline curve equations were applied to 145 gassy coal mines estimated to have closed between 1920 and 1971 in the United States, representing 78 percent of the emissions. State-specific, initial emission rates were used based on average coal mine methane emissions rates during the 1970s (EPA 2003).

Abandoned mines emission estimates are based on all closed mines known to have active mine CH₄ ventilation emission rates greater than 100 mcf/d at the time of abandonment. For example, for 1990 the analysis included 145 mines closed before 1972 and 259 mines closed between 1972 and 1990. Initial emission rates based on MSHA reports, time of abandonment, and basin-specific decline curves influenced by a number of factors were used to calculate annual emissions for each mine in the database. Coal mine degasification data are not available for years prior to 1990, thus the initial emission rates used reflect ventilation emissions only for pre-1990 closures. Methane degasification amounts were added to ventilation data for the total CH₄ liberation rate for fourteen mines that closed between 1992 and 2003. Since the sample of gassy mines (with active mine emissions greater than 100 mcf/d) is assumed to account for 78 percent of the pre-1971 and 98 percent of the post-1971 abandoned mine emissions, the modeled results were multiplied by 1.22 and 1.02 to account for all U.S. abandoned mine emissions. Once the 1991 through 2003 totals were calculated, they were downwardly adjusted to reflect abandoned mine CH₄ emissions avoided from those mines. The inventory totals were not adjusted for abandoned mine reductions in 1990 through 1992, because no data was reported for abandoned coal mining methane recovery projects during that time.

Uncertainty

The parameters for which values must be estimated for each mine in order to predict its decline curve are: 1) the coal's adsorption isotherm; 2) CH₄ flow capacity as expressed by permeability; and 3) pressure at abandonment. Because these parameters are not available for each mine, an approach was used that generates a probability distribution of potential outcomes based on the most likely value and the probable range of values for each parameter. The range of values is not meant to capture the extreme values, but values that represent the highest and lowest quartile of the cumulative probability density function of the parameter. Once the low, mid, and high values are selected, they are applied to a probability density function.

The emission estimates from underground ventilation systems were based on actual measurement data, which are believed to have relatively low uncertainty. A degree of imprecision was introduced because the measurements were not continuous, but rather an average of quarterly instantaneous readings. Additionally, the measurement equipment used possibly resulted in an average of 10 percent overestimation of annual CH₄ emissions (Mutmansky and Wang 2000). Estimates of CH₄ liberated and recovered by degasification systems are also relatively certain because many coal mine operators provided information on individual well gas sales and mined through dates.

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-36. Abandoned coal mines CH₄ emissions in 2003 were estimated to be between 5.4 and 7.8 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 16 percent below to 22 percent above the 2003 emission estimate of 6.4 Tg CO₂ Eq. One of the reasons for the relatively narrow range is that mine-specific data is used in the methodology. The largest degree of uncertainty is associated with the unknown status mines (which account for 41 percent of the mines), with a ±50 percent uncertainty.

Table 3-36: Tier 2 Quantitative Uncertainty Estimates for CH₄ Emissions from Abandoned Underground Coal Mines (Tg CO₂ Eq. and Percent)

Source	Gas	2003	Uncertainty Range Relative to Emission Estimate ^a
		Emission Estimate (Tg CO ₂ Eq.)	
			(%)

			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Abandoned Coal Mines	CH ₄	6.4	5.4	7.8	-16%	+22%

^aRange of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

QA/QC and Verification

As part of a Tier 2 analysis, the United States undertook an effort to verify the model results used in the U.S. Inventory with field measurements. Field measurements were used to test the accuracy of the mathematical decline curves to be used for basin-specific emissions estimates. A series of field measurements were conducted at abandoned mine vent locations across the United States. Between November 1998 and February 2000, EPA recorded measurements at five mines that were not flooded. Measurements were recorded at two abandoned mines located in Ohio and Virginia continuously for 6 to 12 hours. As the methodology was finalized, EPA measured emissions from three additional mines located in Illinois and Colorado. These measurements were recorded hourly for 3 to 4 days and were normalized to average barometric pressures. Prior to these measurements, EPA's Office of Research and Development initiated a field research program in the early 1990s. Data for 21 abandoned mines located throughout the Northern and Central Appalachian, Black Warrior, and Illinois Basins were collected using similar techniques.

Measurements for all field data recorded were plotted against predicted emissions as part of the two studies from 1991 through 2000. Emission rates from nine of the ten mines that were measured fall very close to the predicted mid-case decline rate for their respective basins. For the exponential decline curve fit to the flooding mines, six of nine measurements fall within a 95 percent predictive confidence interval of the mean.

Of the abandoned mines in the database, only about 13 percent of the mines maintain vents to the atmosphere. Therefore, it is difficult to obtain field data. Additional field measurements, however, would be beneficial to further calibrate the equations defined above. Furthermore, it would be useful to extend measurements of diffuse emissions from sealed mines, since they comprise 43 percent of total mines.

Recalculations Discussion

In 2003, all methane emissions from abandoned coal mines were recalculated using: 1) an updated mine list based on 1985 and 1988 USBM records (adding 41 mines to the inventory), 2) updated coal seam permeabilities based on a 2004 EPA study, and 3) revised closure dates for 43 of the mines based on MSHA's data retrieval system. The combination of these changes resulted in an average annual increase of 2.8 Tg CO₂ Eq. (64.5 percent) in CH₄ emissions from abandoned coal mines for the period 1990 through 2002.

3.7. Petroleum Systems (IPCC Source Category 1B2a)

Methane emissions from petroleum systems are primarily associated with crude oil production, transportation, and refining operations. During each of these activities, CH₄ is released to the atmosphere as fugitive emissions, vented emissions, emissions from operational upsets, and emissions from fuel combustion. Total CH₄ emissions from petroleum systems in 2003 were 17.1 Tg CO₂ Eq. (815 Gg). Since 1990, emissions declined due to a decline in domestic oil production and industry efforts to make emissions reductions (see Table 3-37 and Table 3-38). The various sources of emissions are detailed below.

Production Field Operations. Production field operations account for over 95 percent of total CH₄ emissions from petroleum systems. Vented CH₄ from field operations account for approximately 83 percent of the emissions from the production sector, fugitive emissions account for six percent, combustion emissions ten percent, and process upset emissions barely one percent. The most dominant sources of vented emissions are field storage tanks, natural-gas-powered pneumatic devices (low bleed, high bleed, and chemical injection pumps). These four sources alone emit 79 percent of the production field operations emissions. Emissions from storage tanks occur when the CH₄ entrained in crude oil under pressure volatilizes once the crude oil is put into storage tanks at atmospheric pressure. Emissions from high and low-bleed pneumatics occur when pressurized gas that is used for control devices is bled

to the atmosphere as they cycle up and down to modulate the system. Emissions from chemical injection pumps occur as high-pressure gas that is used to drive the pumps is vented to the atmosphere. Two additional large sources, oil well heads and gas engines, together account for 12 percent of emissions from the production sector. The remaining nine percent of the emissions are distributed among 33 additional activities within these four categories.

Crude Oil Transportation. Crude oil transportation activities account for less than one percent of total CH₄ emissions from the oil industry. Venting from tanks and marine vessel loading operations accounts for 65 percent of CH₄ emissions from crude oil transportation. Fugitive emissions, almost entirely from floating roof tanks, account for 18 percent. The remaining 17 percent is distributed among 4 additional sources within these two categories.

Crude Oil Refining. Crude oil refining processes and systems account for only three percent of total CH₄ emissions from the oil industry because most of the CH₄ in crude oil is removed or escapes before the crude oil is delivered to the refineries. There is an insignificant amount of methane in all refined products. Within refineries, vented emissions account for about 87 percent of the emissions, while fugitive and combustion emissions account for approximately six and seven percent respectively. Refinery system blowdowns for maintenance and the process of asphalt blowing—with air, to harden the asphalt—are the primary venting contributors. Most of the fugitive emissions from refineries are from leaks in the fuel gas system. Refinery combustion emissions include small amounts of unburned CH₄ in process heater stack emissions and from unburned CH₄ in engine exhausts and flares.

Table 3-37: CH₄ Emissions from Petroleum Systems (Tg CO₂ Eq.)

Activity	1990	1997	1998	1999	2000	2001	2002	2003
Production Field Operations	19.3	18.1	17.8	17.1	16.9	16.8	16.4	16.4
Pneumatic device venting	11.5	10.8	10.6	10.3	10.0	10.0	9.8	9.8
Tank Venting	3.8	3.4	3.4	3.2	3.2	3.2	3.2	3.2
Combustion & process upsets	2.2	2.0	2.0	1.9	1.9	1.9	1.8	1.8
Misc. venting & fugitives	1.4	1.3	1.3	1.3	1.2	1.2	1.2	1.2
Wellhead fugitives	0.5	0.5	0.5	0.5	0.5	0.5	0.4	0.4
Crude Oil Transportation	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Refining	0.5	0.6	0.6	0.6	0.6	0.6	0.6	0.6
Total estimated emissions	20.0	18.8	18.5	17.8	17.6	17.4	17.1	17.1

Table 3-38: CH₄ Emissions from Petroleum Systems (Gg)

Activity	1990	1997	1998	1999	2000	2001	2002	2003
Production Field Operations	920	862	846	815	803	799	783	782
Pneumatic device venting	545	515	504	488	478	475	465	465
Tank Venting	179	164	162	153	154	154	151	151
Combustion & process upsets	103	96	94	90	89	89	87	86
Misc. venting & fugitives	66	62	61	60	59	59	58	57
Wellhead fugitives	26	25	25	24	22	22	21	21
Crude Oil Transportation	7	6	6	6	5	5	5	5
Refining	25	27	27	27	28	27	27	27
Total estimated emissions	951	895	879	848	836	831	815	815

Methodology

The methodology for estimating CH₄ emissions from petroleum systems is a bottom-up approach, based on a comprehensive study of CH₄ emissions from U.S. petroleum systems (EPA 1999, Radian 1996e). These studies combined emission estimates from 70 activities occurring in petroleum systems from the oil wellhead through crude oil refining, including 39 activities for crude oil production field operations, 11 for crude oil transportation activities, and 20 for refining operations. Annex 3.5 provides greater detail on the emission estimates for these 70

activities. The estimates of CH₄ emissions from petroleum systems do not include emissions downstream of oil refineries because these emissions are very small compared to CH₄ emissions upstream of oil refineries.

The methodology for estimating CH₄ emissions from the 70 oil industry activities employs emission factors initially developed by EPA (1999) and activity factors that are based on EPA (1999) and Radian (1996e) studies. Emissions are estimated for each activity by multiplying emission factors (e.g., emission rate per equipment item or per activity) by their corresponding activity factor (e.g., equipment count or frequency of activity). The report provides emission factors and activity factors for all activities except those related to offshore oil production. For offshore oil production, an emission factor was calculated by dividing an emission estimate from the Minerals Management Service (MMS) by the number of platforms (MMS 2004b). Emission factors were held constant for the period 1990 through 2003.

Activity factors for years 1990 through 2003 were collected from a wide variety of statistical resources. For some years, complete activity factor data were not available. In such cases, one of three approaches was employed. Where appropriate, the activity factor was calculated from related statistics using ratios developed for Radian (1996e). For example, Radian (1996e) found that the number of heater treaters (a source of CH₄ emissions) is related to both number of producing wells and annual production. To estimate the activity factor for heater treaters, reported statistics for wells and production were used, along with the ratios developed for Radian (1996e). In other cases, the activity factor was held constant from 1990 through 2003 based on EPA (1999). Lastly, the previous year's data were used when data for the current year were unavailable. See Annex 3.5 for additional detail.

Nearly all emission factors were taken from Radian (1996e) and EPA (1999). The remaining emission factors were taken from the following sources: EPA default values, MMS reports (MMS 1995), the Exploration and Production (E&P) Tank model (DB Robinson Research Ltd. 1997), and the consensus of industry peer review panels.

Among the more important references used to obtain activity factors are the Energy Information Administration annual and monthly reports (EIA 1990-2003, 1995-2003a-b), the *API Basic Petroleum Data Book* (API 2003), *Methane Emissions from the Natural Gas Industry* by the Gas Research Institute and EPA (Radian 1996a-d), consensus of industry peer review panels, MMS reports (MMS 1995, 2000, 2004a-b), and the *Oil & Gas Journal* (OGJ 2003a-b). Forecasts of activity factors for petroleum systems were developed using production and refining capacity data from the *EIA Annual Energy Outlook* (EIA 2004) as well as offshore activity projections from MMS (2001 and 2004a). Annex 3.5 provides a complete list of references.

Uncertainty

The detailed, bottom-up inventory analysis used to evaluate U.S. petroleum systems reduces the uncertainty related to the CH₄ emission estimates in comparison with a top-down approach. However, some uncertainty still remains. Emission factors and activity factors are based on a combination of measurements, equipment design data, engineering calculations and studies, surveys of selected facilities and statistical reporting. Statistical uncertainties arise from natural variation in measurements, equipment types, operational variability and survey and statistical methodologies. Published activity factors are not available every year for all 70 activities analyzed for petroleum systems; therefore, some are estimated. Because of the dominance of six major sources, which account for 90 percent of the total emissions, the uncertainty surrounding these six sources has been estimated most rigorously, and serves as the basis for determining the overall uncertainty of petroleum systems emission estimates.

The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-39. Petroleum systems CH₄ emissions in 2003 were estimated to be between 11.9 and 51.4 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 30 percent below to 200 percent above the 2003 emission estimate of 17.1 Tg CO₂ Eq.

Table 3-39: Tier 2 Quantitative Uncertainty Estimates for CH₄ Emissions from Petroleum Systems (Tg CO₂ Eq. and Percent)

Source	Gas	2003 Emission	Uncertainty Range Relative to Emission Estimate ^a	
		Estimate (Tg CO ₂ Eq.)	(Tg CO ₂ Eq.)	(%)

			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Petroleum Systems	CH ₄	17.1	11.9	51.4	-30%	+200%

^a Range of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

Recalculations Discussion

Estimates of CH₄ from petroleum systems contain two changes with respect to previous inventories. First, the emission factor for CH₄ emissions from oil tanks in the production sector was modified to remove venting from condensate tanks and only account for venting from crude oil tanks. The previous methodology included an emissions factor that was averaged from the API E&P Tank Calc runs on both oil and condensate tanks. The new calculation is averaged from API E&P Tank Calc runs (DB Robinson Research Ltd. 1997) for API gravity of 44 degrees and below. The adjustment has been made so that vented emissions from condensate tanks can be moved into the emissions from natural gas systems, where they are more relevant. The second change was the development of a new activity factor for offshore Gulf of Mexico platform venting in the production sector. Previously, the activity factor was obtained through MMS, the total number of Gulf of Mexico platforms, and an assumption of what percentage were oil producing platforms. The number of Gulf of Mexico platforms, platform data, and field data is now available on an annual basis from MMS (2000, 2004b, 2004c). From the field and platform data, a new estimate was developed for the percentage of oil producing platforms. This change results in the model reflecting the trend towards more natural gas production than oil production in the Gulf of Mexico. A change in the Gulf of Mexico platform activity resulted in an indirect change in the emissions factor that is calculated from the known base-year emissions value.

The combination of these changes resulted in an average annual decrease of 7.1 Tg CO₂ Eq. (27.4 percent) in CH₄ emissions from petroleum systems for the period 1990 through 2002. Oil tank venting accounted for virtually all of the decrease while Gulf of Mexico platform venting and fugitives remained relatively unchanged.

Planned Improvements

Several improvements to the emission estimates are being evaluated that fine-tune and better track changes in emissions. These include, but are not limited to, some activity factors that are also accounted for in the Natural Gas STAR Program emission reductions, some emission factors for consistency between emission estimates from Petroleum Systems and Natural Gas Systems, and new data from recent studies that bear on both emission factors and activity factors. The growing body of data in the Natural Gas STAR Program, coupled with an increasing number of oil and gas companies doing internal greenhouse gas emissions inventories, provides an opportunity to reevaluate emission and activity factors, as well as the methodology currently used to project emissions from the base year.

3.8. Natural Gas Systems (IPCC Source Category 1B2b)

The U.S. natural gas system encompasses hundreds of thousands of wells, hundreds of processing facilities, and over a million miles of transmission and distribution pipelines. Overall, natural gas systems emitted 125.9 Tg CO₂ Eq. (5,998 Gg) of CH₄ in 2003, a slight decrease over 1990 emissions (see Table 3-40 and Table 3-41). Improvements in management practices and technology, along with the replacement of older equipment, have helped to stabilize emissions (EPA 2002).

Methane emissions from natural gas systems are generally process related, with normal operations, routine maintenance, and system upsets being the primary contributors. Emissions from normal operations include: natural gas combusting engines and turbine exhaust, bleed and discharge emissions from pneumatic devices, and fugitive emissions from system components. Routine maintenance emissions originate from pipelines, equipment, and wells during repair and maintenance activities. Pressure surge relief systems and accidents can lead to system upset emissions. Below is a characterization of the four major stages of the natural gas system. Each of the stages is described and the different factors affecting CH₄ emissions are discussed.

Field Production. In this initial stage, wells are used to withdraw raw gas from underground formations. Emissions arise from the wells themselves, gathering pipelines, and well-site gas treatment facilities such as dehydrators and separators. Fugitive emissions and emissions from pneumatic devices account for the majority of emissions. Emissions from field production accounted for approximately 34 percent of CH₄ emissions from natural gas systems in 2003.

Processing. In this stage, natural gas liquids and various other constituents from the raw gas are removed, resulting in “pipeline quality” gas, which is injected into the transmission system. Fugitive emissions from compressors, including compressor seals, are the primary emission source from this stage. Processing plants account for about 12 percent of CH₄ emissions from natural gas systems.

Transmission and Storage. Natural gas transmission involves high pressure, large diameter pipelines that transport gas long distances from field production and processing areas to distribution systems or large volume customers such as power plants or chemical plants. Compressor station facilities, which contain large reciprocating and turbine compressors, are used to move the gas throughout the United States transmission system. Fugitive emissions from these compressor stations and from metering and regulating stations account for the majority of the emissions from this stage. Pneumatic devices and engine exhaust are also sources of emissions from transmission facilities.

Natural gas is also injected and stored in underground formations, or liquefied and stored in above ground tanks, during periods of low demand (e.g., summer), and withdrawn, processed, and distributed during periods of high demand (e.g., winter). Compressors and dehydrators are the primary contributors to emissions from these storage facilities. Methane emissions from transmission and storage sector account for approximately 32 percent of emissions from natural gas systems.

Distribution. Distribution pipelines take the high-pressure gas from the transmission system at “city gate” stations, reduce the pressure and distribute the gas through primarily underground mains and service lines to individual end users. There were over 978,000 miles of distribution mains in 2003, an increase from just over 789,500 miles in 1990 (OPS 2002a). Distribution system emissions, which account for approximately 22 percent of emissions from natural gas systems, result mainly from fugitive emissions from gate stations and non-plastic piping (cast iron, steel).⁵¹ An increased use of plastic piping, which has lower emissions than other pipe materials, has reduced the growth in emissions from this stage. Distribution system emissions in 2003 were 10 percent lower than 1990 levels.

Table 3-40: CH₄ Emissions from Natural Gas Systems (Tg CO₂ Eq.)*

Stage	1990	1997	1998	1999	2000	2001	2002	2003
Field Production	36.3	40.4	41.2	37.5	41.0	44.5	44.5	43.4
Processing	14.8	14.9	14.7	14.6	14.9	15.1	14.6	14.5
Transmission and Storage	46.8	46.1	44.7	43.4	43.4	40.2	42.0	40.6
Distribution	30.5	32.4	31.2	31.9	32.8	32.1	29.5	27.4
Total	128.3	133.6	131.8	127.4	132.1	131.8	130.6	125.9

*Including CH₄ emission reductions achieved by the Natural Gas STAR program.

Note: Totals may not sum due to independent rounding.

Table 3-41: CH₄ Emissions from Natural Gas Systems (Gg)*

Stage	1990	1997	1998	1999	2000	2001	2002	2003
Field Production	1,731	1,921	1,962	1,785	1,953	2,117	2,121	2,068
Processing	704	708	702	696	708	717	693	691
Transmission and Storage	2,226	2,193	2,127	2,068	2,067	1,914	2,002	1,933
Distribution	1,450	1,541	1,484	1,517	1,560	1,530	1,405	1,305

⁵¹ The percentages of total emissions from each stage may not add to 100 because of independent rounding.

Total	6,112	6,363	6,276	6,066	6,289	6,277	6,221	5,998
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*Including CH₄ emission reductions achieved by the Natural Gas STAR program.

Note: Totals may not sum due to independent rounding.

Methodology

The basis for estimates of CH₄ emissions from the U.S. natural gas industry is a detailed study by the Gas Research Institute and EPA (EPA/GRI 1996). The EPA/GRI study developed over 100 emission and activity factors to characterize emissions from the various components within the operating stages of the U.S. natural gas system. The study was based on a combination of process engineering studies and measurements at representative gas facilities. From this analysis, a 1992 emission estimate was developed using the emission and activity factors. For other years, a set of industry activity factor drivers was developed that can be used to update activity factors. These drivers include statistics on gas production, number of wells, system throughput, miles of various kinds of pipe, and other statistics that characterize the changes in the U.S. natural gas system infrastructure and operations.

See Annex 3.4 for more detailed information on the methodology and data used to calculate CH₄ emissions from natural gas systems.

Activity factor data were taken from the following sources: American Gas Association (AGA 1991-1998); American Petroleum Institute (API 2002, 2003); Annual Energy Review (EIA 2002f); Historical Natural Gas Annual (EIA 2003e); Minerals and Management Service (MMS 1998, 1999, 2000, 2001, 2002a-b, 2003, 2004a-d); Monthly Energy Review (EIA 2004d); Natural Gas Annual (EIA 1993, 1996a, 1997a, 1998a-b, 2001a, 2002a, 2003a); Natural Gas Liquids Reserves Report (EIA 1996b, 1997b, 1998c, 1999, 2000, 2001c, 2002d); Natural Gas Monthly (EIA 2001b, 2002b-c, 2003b-d, 2004a-c); the Natural Gas STAR Program annual emissions savings (EPA 2004); Oil and Gas Journal (OGJ 1999 - 2004); Office of Pipeline Safety (OPS 2004a-b) other Energy Information Administration publications (EIA 2002e, 2004e-f). The Gas Systems Analysis model was used to aid in collecting data for non-associated and associated wells (GSAM 1997). Data from a program for estimating emissions from hydrocarbon production tanks is incorporated (DB Robinson Research Ltd. 1997). Coalbed Methane well activity factors were taken from the Wyoming Oil and Gas Conservation Commission (Wyoming 2004) and the Alabama State Oil and Gas Board (Alabama 2004). Other state well data was taken from : American Association of Petroleum Geologists (AAPG 2204); Brookhaven College (Brookhaven 2004); Kansas Geological Survey (Kansas 2004); Rocky Mountain Production Report (Lippman (2003); Montana Board of Oil and Gas Conservation (Montana 2004); Oklahoma Geological Survey (Oklahoma 2004); Utah Division of Oil, Gas and Mining (Utah 2004). Emissions factors were taken from EPA/GRI (1996).

Uncertainty

The heterogeneous nature of the natural gas industry makes it difficult to sample facilities that are completely representative of the entire industry. Because of this, scaling up from model facilities introduces a degree of uncertainty. Additionally, highly variable emission rates were measured among many system components, making the calculated average emission rates uncertain. The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-42. Natural gas systems CH₄ emissions in 2003 were estimated to be between 87.1 and 166.7 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 31 percent below to 32 percent above the 2003 emission estimate of 125.9 Tg CO₂ Eq.

Table 3-42: Tier 2 Quantitative Uncertainty Estimates for CH₄ Emissions from Natural Gas Systems (Tg CO₂ Eq. and Percent)

Source	Gas	2003	Uncertainty Range Relative to Emission			
		Emission Estimate	Estimate ^a			
		(Tg CO ₂ Eq.)	(Tg CO ₂ Eq.)	(%)		
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Natural Gas	CH ₄	125.9	87.1	166.7	-31%	+32%

Recalculations Discussion

Emissions with Natural Gas STAR Program reductions were updated using new Gas STAR emissions reduction data from the iSTAR database. Gas STAR reductions reported retroactively for the years 1990 to 1992 are assumed to be characterized in the GRI/EPA 1996 study for the base year 1992. Therefore, to avoid double counting, those emissions reductions are not counted in the inventory. Gas STAR reductions are also subject to sunseting rules that prevent perpetual crediting of all reductions. The inventory assumes that once an emissions reducing technology is put in place, it will continue to provide emissions savings and thus the sunseting rule is not implemented in the inventory.

Three new sources of emissions in the production sector of the natural gas systems were added this year:

- Gas condensate stored in tanks vents methane and other hydrocarbons to the atmosphere. Emissions from condensate tanks can be divided into two categories: tanks with control devices such as vapor recovery units or flares, and tanks that have no control devices. Condensate tanks contributed an estimated 52.3 Gg of methane emissions that represents 0.87 percent of total methane emissions from natural gas systems in 2003. In 1990, Condensate tanks were estimated to contribute 39.9 Gg of emissions or about 0.65 percent of total emissions from natural gas systems.
- Another source added to the production sector for the 2003 reporting year was unconventional gas well fugitives. Wells in this source were previously treated as conventional wells in the rest of U.S. (western) geographic region. Unconventional well fugitives include fugitive emissions from coal bed CH₄ and shale wells. From 1990 through 2003, unconventional well fugitives accounted for a small amount of emissions from natural gas systems.
- The production sector also had flaring from offshore Gulf of Mexico operations added, using a 2 percent uncombusted hydrocarbon factor. This source contributed to less than 0.1 percent to the emissions from the entire time series. A second Gulf of Mexico source added to the production sector was Gulf of Mexico offshore well venting. This contributed about 2.4 percent of the 2003 emissions and about 2.6 percent in 1990.

The combination of these methodological and historical data changes resulted in an average annual increase of 6.7 Tg CO₂ Eq. (5.3 percent) in CH₄ emissions from natural gas systems for the period 1990 through 2002.

Planned Improvements

Several improvements to the emission estimates are being evaluated that fine-tune and better track changes in emissions. These include, but are not limited to, some activity factors that are also accounted for in the Natural Gas STAR Program emission reductions, some emission factors for consistency between emission estimates from the Petroleum Systems and Natural Gas Systems source categories, and new data from recent studies that bear on both emission factors and activity factors. The growing body of data in the Natural Gas STAR Program, coupled with an increasing number of oil and gas companies doing internal greenhouse gas emissions inventories, provides an opportunity to reevaluate emission and activity factors, as well as the methodology currently used to project emissions from the base year. Two improvements that are of particular note are in the production and processing sector. Recent data has suggested that the emission factor for well clean-ups in the production sector is low, and additional data sources are being investigated to produce a more robust emission factor. In the processing sector, improvements are being considered across the entire sector based on studies completed and near completion at five or more processing plants. These studies suggest the need for changing several emission factors as well as adding some additional sources within the processing sector.

3.9. Municipal Solid Waste Combustion (IPCC Source Category 1A5)

Combustion is used to manage about 7 to 17 percent of the municipal solid wastes generated in the United States, depending on the source of the estimate and the scope of materials included in the definition of solid waste (EPA 2000c, Goldstein and Matdes 2001, Kaufman et al. 2004). Almost all combustion of municipal solid wastes in the United States occurs at waste-to-energy facilities where energy is recovered, and thus emissions from waste combustion are accounted for in the Energy chapter. Combustion of municipal solid wastes results in conversion of the organic inputs to CO₂. According to the IPCC Guidelines, when the CO₂ emitted is of fossil origin, it is counted as a net anthropogenic emission of CO₂ to the atmosphere. Thus, the emissions from waste combustion are calculated by estimating the quantity of waste combusted and the fraction of the waste that is carbon derived from fossil sources.

Most of the organic materials in municipal solid wastes are of biogenic origin (e.g., paper, yard trimmings), and have their net carbon flows accounted for under the Land-Use Change and Forestry chapter (see Box 3-3). However, some components—plastics, synthetic rubber, synthetic fibers, and carbon black—are of fossil origin. Plastics in the U.S. waste stream are primarily in the form of containers, packaging, and durable goods. Rubber is found in durable goods, such as carpets, and in non-durable goods, such as clothing and footwear. Fibers in municipal solid wastes are predominantly from clothing and home furnishings. Tires (which contain rubber and carbon black) are also considered a “non-hazardous” waste and are included in the municipal solid waste combustion estimate, though waste disposal practices for tires differ from the rest of municipal solid waste.

[Begin Text Box]

Box 3-3: Biogenic Emissions and Sinks of Carbon

For many countries, CO₂ emissions from the combustion or degradation of biogenic materials are important because of the significant amount of energy they derive from biomass (e.g., burning fuelwood). The fate of biogenic materials is also important when evaluating waste management emissions (e.g., the decomposition of paper). The carbon contained in paper was originally stored in trees during photosynthesis. Under natural conditions, this material would eventually degrade and cycle back to the atmosphere as CO₂. The quantity of carbon that these degradation processes cycle through the Earth’s atmosphere, waters, soils, and biota is much greater than the quantity added by anthropogenic greenhouse gas sources. However, the focus of the UNFCCC is on emissions resulting from human activities and subject to human control, because it is these emissions that have the potential to alter the climate by disrupting the natural balances in carbon’s biogeochemical cycle, and enhancing the atmosphere’s natural greenhouse effect.

Carbon dioxide emissions from the combustion or decomposition of biogenic materials (e.g., paper, wood products, and yard trimmings) grown on a sustainable basis are considered to mimic the closed loop of the natural carbon cycle—that is, they return to the atmosphere CO₂ that was originally removed by photosynthesis. However, CH₄ emissions from landfilled waste occur due to the man-made anaerobic conditions conducive to CH₄ formation that exist in landfills, and are consequently included in this inventory.

The removal of carbon from the natural cycling of carbon between the atmosphere and biogenic materials—which occurs when wastes of biogenic origin are deposited in landfills—sequesters carbon. When wastes of sustainable, biogenic origin are landfilled, and do not completely decompose, the carbon that remains is effectively removed from the global carbon cycle. Landfilling of forest products, yard trimmings, and food scraps resulted in net long-term storage of 10.1 Tg CO₂ Eq. in 2003, as described in the Land-Use Change and Forestry chapter.

[End Box]

Approximately 24 million metric tons of municipal solid wastes were combusted in the United States in 2003. Carbon dioxide emissions from combustion of municipal solid wastes rose 72 percent since 1990, to an estimated 18.8 Tg CO₂ Eq. (18,781 Gg) in 2003, as the volume of plastics and other fossil carbon-containing materials in MSW increased (see Table 3-43 and Table 3-44). Waste combustion is also a source of N₂O emissions (De Soete

1993). Nitrous oxide emissions from municipal solid waste combustion were estimated to be 0.5 Tg CO₂ Eq. (1 Gg) in 2003, and have not changed significantly since 1990.

Table 3-43: CO₂ and N₂O Emissions from Municipal Solid Waste Combustion (Tg CO₂ Eq.)

Gas/Waste Product	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	10.9	17.8	17.1	17.6	18.0	18.8	18.8	18.8
Plastics	8.0	11.9	11.4	12.0	12.1	12.7	12.7	12.7
Synthetic Rubber in Tires	0.2	0.9	0.9	0.9	0.9	0.9	0.9	0.9
Carbon Black in Tires	0.2	1.2	1.2	1.2	1.2	1.2	1.2	1.2
Synthetic Rubber in MSW	1.3	1.7	1.6	1.6	1.7	1.8	1.8	1.8
Synthetic Fibers	1.2	2.1	2.0	2.0	2.1	2.2	2.2	2.2
N₂O	0.4	0.4	0.3	0.3	0.4	0.5	0.5	0.5
Total	11.3	18.1	17.4	18.0	18.3	19.2	19.2	19.2

Table 3-44: CO₂ and N₂O Emissions from Municipal Solid Waste Combustion (Gg)

Gas/Waste Product	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	10,919	17,761	17,094	17,632	17,979	18,781	18,781	18,781
Plastics	7,953	11,914	11,427	11,950	12,145	12,718	12,718	12,718
Synthetic Rubber in Tires	191	891	887	890	893	895	895	895
Carbon Black in Tires	249	1,165	1,160	1,164	1,167	1,170	1,170	1,170
Synthetic Rubber in MSW	1,330	1,725	1,627	1,612	1,689	1,810	1,810	1,810
Synthetic Fibers	1,196	2,065	1,992	2,016	2,086	2,187	2,187	2,187
N₂O	1	1	1	1	1	1	1	1

Ambient air pollutants are also emitted during waste incineration and open burning, as shown in Table 3-45. These emissions are a relatively small portion of the overall ambient air pollutant emissions, comprising less than 5 percent for each gas over the entire time series.

Table 3-45: NO_x, CO, and NMVOC Emissions from Municipal Solid Waste Combustion (Gg)

Gas/Source	1990	1997	1998	1999	2000	2001	2002	2003
NO_x	82	140	145	143	114	114	134	121
Waste Incineration	44	48	49	48	38	38	45	41
Open Burning	38	92	96	95	76	76	89	80
CO	978	2,668	2,826	2,725	1,670	1,672	1,672	1,674
Waste Incineration	337	68	69	66	40	41	41	41
Open Burning	641	2,600	2,757	2,659	1,630	1,631	1,631	1,633
NMVOCs	222	313	326	302	257	258	281	263
Waste Incineration	44	23	23	19	15	16	18	16
Open Burning	178	290	303	284	242	242	264	246

Note: Totals may not sum due to independent rounding.

Methodology

Emissions of CO₂ from MSW combustion include CO₂ generated by the combustion of plastics, synthetic fibers, and synthetic rubber, as well as the combustion of synthetic rubber and carbon black in tires. These emissions were calculated by multiplying the amount of each material combusted by the carbon content of the material and the fraction oxidized (98 percent). Plastics combusted in municipal solid wastes were categorized into seven plastic resin types, each material having a discrete carbon content. Similarly, synthetic rubber is categorized into three product types, and synthetic fibers were categorized into four product types, each having a discrete carbon content. Scrap tires contain several types of synthetic rubber, as well as carbon black. Each type of synthetic rubber has a discrete carbon content, and carbon black is 100 percent carbon. Emissions of CO₂ were calculated based on the

number of scrap tires used for fuel and the synthetic rubber and carbon black content of the tires. More detail on the methodology for calculating emissions from each of these waste combustion sources is provided in Annex 3.6.

For each of the methods used to calculate CO₂ emissions from municipal solid waste combustion, data on the quantity of product combusted and the carbon content of the product are needed. For plastics, synthetic rubber, and synthetic fibers, the amount of material in municipal solid wastes and its portion combusted were taken from the *Characterization of Municipal Solid Waste in the United States* (EPA 2000c, 2002a, 2003). For synthetic rubber and carbon black in scrap tires, this information was provided by the *U.S. Scrap Tire Markets 2001* (RMA 2002) and *Scrap Tires, Facts and Figures* (STMC 2000, 2001, 2002, 2003). Data were not available for 2002 or 2003, so the values for these years were assumed to equal the value for 2001.

Average carbon contents for the “Other” plastics category, synthetic rubber in municipal solid wastes, and synthetic fibers were calculated from 1998 production statistics, which divide their respective markets by chemical compound. For synthetic rubber in scrap tires information about scrap tire composition was taken from the Scrap Tire Management Council’s internet site (STMC 2003).

The assumption that 98 percent of organic carbon is oxidized (which applies to all municipal solid waste combustion categories for CO₂ emissions) was reported in the EPA’s life cycle analysis of greenhouse gas emissions and sinks from management of solid waste (EPA 2002b).

Combustion of municipal solid waste also results in emissions of N₂O. These emissions were calculated as a function of the total estimated mass of municipal solid waste combusted and an emission factor. The N₂O emission estimates are based on different data sources. As noted above, N₂O emissions are a function of total waste combusted in each year; for 1990 through 2002, these data were derived from the information published in *BioCycle* (Kaufman et al 2004). As for the activity data for CO₂ emissions, data on total waste combusted was not available for 2003, so the value for this year was assumed to equal the most recent value available (2002). Table 3-46 provides data on municipal solid waste generation and percentage combustion for the total waste stream. The emission factor of N₂O emissions per quantity of municipal solid waste combusted is an average of values from IPCC’s *Good Practice Guidance* (2000).

Table 3-46: Municipal Solid Waste Generation (Metric Tons) and Percent Combusted

Year	Waste Generation	Combusted (%)
1990	266,365,714	11.5
1991	254,628,360	10.0
1992	264,668,342	11.0
1993	278,388,835	10.0
1994	292,915,829	10.0
1995	296,390,405	10.0
1996	297,071,712	10.0
1997	308,870,755	9.0
1998	339,865,243	7.5
1999	347,089,277	7.0
2000	371,071,109	7.0
2001	404,002,786 ^a	7.4 ^a
2002	436,934,464	7.7
2003	436,934,464 ^b	7.7 ^b

^a Interpolated between 2000 and 2002 values.

^b Assumed equal to 2002 value.

EPA (2003) provided emission estimates for NO_x, CO, and NMVOCs from waste incineration and open burning, which were determined using industry published production data and applying average emission factors.

Uncertainty

A Tier 2 Monte Carlo analysis was performed to determine the level of uncertainty surrounding the estimates of CO₂ emissions and N₂O emissions from municipal solid waste combustion. IPCC Tier 2 analysis allows the specification of probability density functions for key variables within a computational structure that mirrors the calculation of the inventory estimate. Uncertainty estimates and distributions for waste generation variables (i.e., plastics, synthetic rubber, and textiles generation) were obtained through a conversation with one of the authors of the *Municipal Solid Waste in the United States* reports. Statistical analyses or expert judgments of uncertainty were not available directly from the information sources for the other variables; thus, uncertainty estimates for these variables were determined using assumptions based on source category knowledge and the known uncertainty estimates for the waste generation variables. The highest levels of uncertainty surround the variables that are based on assumptions (e.g., percent of clothing and footwear composed of synthetic rubber); the lowest levels of uncertainty surround variables that were determined by quantitative measurements (e.g., combustion efficiency, carbon content of carbon black).

The results of the 2003 uncertainty analysis of CO₂ emissions are the same as the 2002 results, given that the data for the two years is identical (no data updated for 2003 were available). The results of the Tier 2 quantitative uncertainty analysis are summarized in Table 3-47. Municipal solid waste combustion CO₂ emissions in 2003 were estimated to be between 15.2 and 21.6 Tg CO₂ Eq. at a 95 percent confidence level (or in 19 out of 20 Monte Carlo Simulations). This indicates a range of 19 percent below to 15 percent above the 2003 emission estimate of 18.8 Tg CO₂ Eq. Also at a 95 percent confidence level, municipal solid waste combustion N₂O emissions in 2003 were estimated to be between 0.13 and 1.34 Tg CO₂ Eq. This indicates a range of 71 percent below to 192 percent above the 2003 emission estimate of 0.5 Tg CO₂ Eq.

Table 3-47: Tier 2 Quantitative Uncertainty Estimates for CO₂ and N₂O from Municipal Solid Waste Combustion (Tg CO₂ Eq. and Percent)

Source	Gas	2003 Emission Estimate (Tg CO ₂ Eq.)	Uncertainty Range Relative to Emission Estimate ^a			
			(Tg CO ₂ Eq.)		(%)	
			Lower Bound	Upper Bound	Lower Bound	Upper Bound
Municipal Solid Waste Combustion	CO ₂	18.8	15.2	21.6	-19%	+15%
Municipal Solid Waste Combustion	N ₂ O	0.5	0.13	1.34	-71%	+192%

^a Range of emission estimates predicted by Monte Carlo Simulation for a 95 percent confidence interval.

The uncertainties in the waste combustion emission estimates arise from both the assumptions applied to the data and from the quality of the data.

- MSW Combustion Rate.** A source of uncertainty affecting both fossil CO₂ and N₂O emissions is the estimate of the MSW combustion rate. The EPA (2000c, 2002a, 2003) estimates of materials generated, discarded, and combusted carry considerable uncertainty associated with the material flows methodology used to generate them. Similarly, the *BioCycle* (Glenn 1999, Goldstein and Matdes 2000, Goldstein and Matdes 2001, Kaufman et al. 2004) estimate of total waste combustion—used for the N₂O emissions estimate—is based on a survey of state officials, who use differing definitions of solid waste and who draw from a variety of sources of varying reliability and accuracy. The survey methodology changed significantly and thus the results reported for 2002 are not directly comparable to the earlier results (Kaufman et al. 2004), introducing further uncertainty. Despite the differences in methodology and data sources, the two references—the EPA’s Office of Solid Waste (EPA 2000a, 2002b, 2003) and the *BioCycle* series—provide estimates of total solid waste combusted that are relatively consistent (see Table 3-48).

Table 3-48: U.S. Municipal Solid Waste Combusted, as Reported by EPA and BioCycle (Metric Tons)

Year	EPA	BioCycle
1990	28,855,809	30,632,057
1991	27,773,783	25,462,836
1992	29,568,442	29,113,518
1993	28,696,188	27,838,884
1994	29,532,844	29,291,583
1995	32,182,194	29,639,040
1996	32,831,450	29,707,171
1997	33,597,844	27,798,368
1998	31,205,358	25,489,893
1999	30,859,134	24,296,249
2000	30,512,946	25,974,978
2001	30,569,746	29,694,205 ^a
2002	NA	33,643,954
2003	NA	NA

NA (Not Available)

^a Interpolated between 2000 and 2002 values.

- Fraction Oxidized.* Another source of uncertainty for the CO₂ emissions estimate is fraction oxidized. Municipal waste combustors vary considerably in their efficiency as a function of waste type, moisture content, combustion conditions, and other factors. Despite this variability in oxidation rates, a value of 98 percent was assumed for this analysis.
- Missing Data on Municipal Solid Waste Composition.* Disposal rates have been interpolated when there is an incomplete interval within a time series. Where data are not available for years at the end of a time series (1990, 2003), they are set equal to the most recent years for which estimates are available.
- Average Carbon Contents.* Average carbon contents were applied to the mass of “Other” plastics combusted, synthetic rubber in tires and municipal solid waste, and synthetic fibers. These average values were estimated from the average carbon content of the known products recently produced. The true carbon content of the combusted waste may differ from this estimate depending on differences in the chemical formulation between the known and unspecified materials, and differences between the composition of the material disposed and that produced. For rubber, this uncertainty is probably small since the major elastomers’ carbon contents range from 77 to 91 percent; for plastics, where carbon contents range from 29 to 92 percent, it may be more significant. Overall, this is a small source of uncertainty.
- Synthetic/Biogenic Assumptions.* A portion of the fiber and rubber in municipal solid waste is biogenic in origin. Assumptions have been made concerning the allocation between synthetic and biogenic materials based primarily on expert judgment.
- Combustion Conditions Affecting N₂O Emissions.* Because insufficient data exist to provide detailed estimates of N₂O emissions for individual combustion facilities, the estimates presented exhibit high uncertainty. The emission factor for N₂O from municipal solid waste combustion facilities used in the analysis is an average of default values used to estimate N₂O emissions from facilities worldwide (Johnke 1999, UK: Environment Agency 1999, Yasuda 1993). These factors span an order of magnitude, reflecting considerable variability in the processes from site to site. Due to a lack of information on the control of N₂O emissions from MSW combustion facilities in the United States, the estimate of zero percent for N₂O emissions control removal efficiency also exhibits uncertainty.

Recalculations Discussion

The N₂O emissions estimates for 2001 and 2002 are slightly different from those reported in last year’s inventory because newly-available data (Kaufman et al. 2004) were used for the tonnage of waste burned in municipal solid

waste combustion for those years. The change resulted in 2001 and 2002 N₂O emissions from municipal solid waste combustion that increased by less than 0.1 Tg CO₂ Eq. (14.7 percent and 26.4 percent, respectively).

The NO_x, CO, and NMVOC emissions estimates for 1999 through 2002 also vary slightly from last year's inventory due to revised criteria pollutant data from EPA. The change constituted less than a 1687 Gg decrease in NO_x, CO, and NMVOC emissions in 2002.

3.10. Natural Gas Flaring and Ambient Air Pollutant Emissions from Oil and Gas Activities (IPCC Source Category 1B2)

The flaring of natural gas from on- and off-shore oil wells is a small source of CO₂. In addition, oil and gas activities also release small amounts of NO_x, CO, and NMVOCs. This source accounts for only a small proportion of overall emissions of each of these gases. Emissions of NO_x and CO from petroleum and natural gas production activities were both less than 1 percent of national totals, while NMVOC and SO₂ emissions were roughly 2 percent of national totals.

The flaring (i.e. combustion) and venting of natural gas during petroleum production result in the release of CO₂ and CH₄ emissions, respectively. Barns and Edmonds (1990) noted that of total reported U.S. venting and flaring, approximately 20 percent may be vented, with the remaining 80 percent flared, but it is now believed that flaring accounts for an even greater proportion. Studies indicate that the percentage of natural gas that is flared from off-shore U.S. production is considerably lower (approximately 30 percent in 2003), due in part to differences in the legislation governing on- and off-shore natural gas production. Methane emissions from venting are accounted for in the Petroleum Systems source category. For 2003, total CO₂ emissions from flaring activities were estimated to be 5.9 Tg CO₂ Eq. (5,970 Gg), an increase of 3 percent from 1990 levels. On-shore flaring activities accounted for 5.7 Tg CO₂ Eq. (5,743 Gg), or 96 percent, of the total flaring emissions, while off-shore flaring constituted 0.2 Tg CO₂ Eq. (227 Gg), or 4 percent, of the total (see Table 3-49).

Table 3-49: CO₂ Emissions from On-Shore and Off-Shore Natural Gas Flaring (Tg CO₂ Eq.)

	1990	1997	1998	1999	2000	2001	2002	2003
On-Shore Flaring	5.5	7.6	6.3	6.7	5.5	5.9	6.0	5.7
Off-Shore Flaring	0.3	0.3	0.3	0.3	0.2	0.2	0.2	0.2
Total Flaring	5.8	7.9	6.6	7.0	5.8	6.1	6.2	5.9

Note: Totals may not sum due to independent rounding.

Table 3-50: CO₂ Emissions from On-Shore and Off-Shore Natural Gas Flaring (Gg)

	1990	1997	1998	1999	2000	2001	2002	2003
On-Shore Flaring	5,514	7,565	6,250	6,679	5,525	5,858	6,006	5,743
Off-Shore Flaring	296	309	316	264	244	236	227	227
Total Flaring	5,810	7,874	6,566	6,943	5,769	6,094	6,233	5,970

Note: Totals may not sum due to independent rounding.

In addition, oil and gas activities, including production, transportation, and storage, result in the release of small amounts of NO_x, CO, and NMVOCs. Ambient air pollutant emissions from this source from 1990 to 2003 are presented below (see Table 3-51).

Table 3-51: NO_x, NMVOCs, and CO Emissions from Oil and Gas Activities (Gg)

Year	NO _x	CO	NMVOCs
1990	139	302	555
1996	126	321	433
1997	130	333	442
1998	130	332	440
1999	109	145	414
2000	111	146	389
2001	113	147	400

2002	135	116	340
2003	124	125	345

Methodology

Estimates of CO₂ emissions from on- and off-shore natural gas flaring were prepared using an emission factor of 54.71 Tg CO₂ Eq./QBtu of flared gas, and an assumed flaring efficiency of 100 percent. Ambient air pollutant emission estimates for NO_x, CO, and NMVOCs were determined using industry-published production data and applying average emission factors.

Total on-shore natural gas vented and flared was taken from EIA's *Natural Gas Annual* (EIA 2004); however, there is a discrepancy in the time series. One facility in Wyoming had been incorrectly reporting CO₂ vented as CH₄. EIA noted and corrected these data in the *Natural Gas Annual 2000* (EIA 2001) for the years 1998 and 1999 only. Data for 1990 through 1997 were adjusted by assuming a proportionate share of CO₂ in the flare gas for those years as for 1998 and 1999. The adjusted values are provided in Table 3-52. It was assumed that all reported vented and flared gas was flared. This assumption is consistent with that used by EIA in preparing their emission estimates, under the assumption that many states require flaring of natural gas (EIA 2000b). The emission and thermal conversion factors were also provided by EIA (2001) and are included in Table 3-52.

The total off-shore natural gas vented and flared was obtained from the Minerals Management Service's OGOR-B reports (MMS 2003). The percentage of natural gas flared was estimated using data from a 1993 air quality study and emissions inventory of the Gulf of Mexico (MOADS) and a 2000 emissions inventory conducted for the Breton National Wilderness Area Management Plan (BOADS). See Table 3-53

Emission estimates for NO_x, CO, and NMVOCs from petroleum refining, petroleum product storage and transfer, and petroleum marketing operations were obtained from preliminary data (EPA 2003), which, in its final iteration, will be published on the National Emission Inventory (NEI) Air Pollutant Emission Trends web site. Included are gasoline, crude oil and distillate fuel oil storage and transfer operations, gasoline bulk terminal and bulk plants operations, and retail gasoline service stations operations.

Table 3-52: Total Natural Gas Reported Vented and Flared (Million Ft³) and Thermal Conversion Factor (Btu/Ft³)

Year	Vented and Flared (original)	Vented and Flared (revised)*	Thermal Conversion Factor
1990	150,415	91,130	1,105
1991	169,909	92,207	1,108
1992	167,519	83,363	1,110
1993	226,743	108,238	1,106
1994	228,336	109,493	1,105
1995	283,739	144,265	1,106
1996	272,117	135,709	1,109
1997	256,351	124,918	1,107
1998	103,019	103,019	1,109
1999	110,285	110,285	1,107
2000	91,232	91,232	1,107
2001	96,913	96,913	1,105
2002	99,173	99,173	1,107
2003	94,929	94,929	1,106

* Wyoming venting and flaring estimates were revised. See text for further explanation.

Table 3-53: Volume Flared Offshore (MMcf) and Fraction Vented and Flared (Percent)

	1990	1997	1998	1999	2000	2001	2002	2003
Total Gulf of Mexico (GOM)								
Vented & Flared (MMcf)	13,610	15,440	16,280	14,057	12,971	12,990	12,487	12,487

Estimated Flaring Fraction of GOM Vented & Flared	36%	33%	32%	31%	31%	30%	30%	30%
Total	4,900	5,095	5,210	4,358	4,021	3,897	3,746	3,746

Uncertainty

Uncertainties in CO₂ emission estimates primarily arise from assumptions concerning the flaring efficiency and the correction factor applied to 1990 through 1997 venting and flaring data. Uncertainties in ambient air pollutant emission estimates are partly due to the accuracy of the emission factors used and projections of growth.

Recalculations Discussion

The historical data for natural gas flaring was adjusted slightly, which resulted in an average annual increase in CO₂ emissions from flaring of 0.1 Tg CO₂ Eq. (2.3 percent) for the period 1990 through 2002.

3.11. International Bunker Fuels (IPCC Source Category 1: Memo Items)

Emissions resulting from the combustion of fuels used for international transport activities, termed international bunker fuels under the UNFCCC, are currently not included in national emission totals, but are reported separately based upon location of fuel sales. The decision to report emissions from international bunker fuels separately, instead of allocating them to a particular country, was made by the Intergovernmental Negotiating Committee in establishing the Framework Convention on Climate Change.⁵² These decisions are reflected in the *Revised 1996 IPCC Guidelines*, in which countries are requested to report emissions from ships or aircraft that depart from their ports with fuel purchased within national boundaries and are engaged in international transport separately from national totals (IPCC/UNEP/OECD/IEA 1997).⁵³

Greenhouse gases emitted from the combustion of international bunker fuels, like other fossil fuels, include CO₂, CH₄, N₂O, CO, NO_x, NMVOCs, particulate matter, and SO₂.⁵⁴ Two transport modes are addressed under the IPCC definition of international bunker fuels: aviation and marine.⁵⁵ Emissions from ground transport activities—by road vehicles and trains—even when crossing international borders are allocated to the country where the fuel was loaded into the vehicle and, therefore, are not counted as bunker fuel emissions.

The IPCC Guidelines distinguish between different modes of air traffic. Civil aviation comprises aircraft used for the commercial transport of passengers and freight, military aviation comprises aircraft under the control of national armed forces, and general aviation applies to recreational and small corporate aircraft. The IPCC Guidelines further define international bunker fuel use from civil aviation as the fuel combusted for civil (e.g., commercial) aviation purposes by aircraft arriving or departing on international flight segments. However, as mentioned above, and in keeping with the IPCC Guidelines, only the fuel purchased in the United States and used by aircraft taking-off (i.e.,

⁵² See report of the Intergovernmental Negotiating Committee for a Framework Convention on Climate Change on the work of its ninth session, held at Geneva from 7 to 18 February 1994 (A/AC.237/55, annex I, para. 1c).

⁵³ Note that the definition of international bunker fuels used by the UNFCCC differs from that used by the International Civil Aviation Organization.

⁵⁴ Sulfur dioxide emissions from jet aircraft and marine vessels, although not estimated here, are mainly determined by the sulfur content of the fuel. In the United States, jet fuel, distillate diesel fuel, and residual fuel oil average sulfur contents of 0.05, 0.3, and 2.3 percent, respectively. These percentages are generally lower than global averages.

⁵⁵ Most emission related international aviation and marine regulations are under the rubric of the International Civil Aviation Organization (ICAO) or the International Maritime Organization (IMO), which develop international codes, recommendations, and conventions, such as the International Convention of the Prevention of Pollution from Ships (MARPOL).

departing) from the United States are reported here. The standard fuel used for civil aviation is kerosene-type jet fuel, while the typical fuel used for general aviation is aviation gasoline.⁵⁶

Emissions of CO₂ from aircraft are essentially a function of fuel use. Methane, N₂O, CO, NO_x, and NMVOC emissions also depend upon engine characteristics, flight conditions, and flight phase (i.e., take-off, climb, cruise, decent, and landing). Methane, CO, and NMVOCs are the product of incomplete combustion and occur mainly during the landing and take-off phases. In jet engines, N₂O and NO_x are primarily produced by the oxidation of atmospheric nitrogen, and the majority of emissions occur during the cruise phase. The impact of NO_x on atmospheric chemistry depends on the altitude of the actual emission. The cruising altitude of supersonic aircraft, near or in the ozone layer, is higher than that of subsonic aircraft. At this higher altitude, NO_x emissions contribute to stratospheric ozone depletion.⁵⁷ At the cruising altitudes of subsonic aircraft, however, NO_x emissions contribute to the formation of tropospheric ozone. At these lower altitudes, the positive radiative forcing effect of ozone has enhanced the anthropogenic greenhouse gas forcing.⁵⁸ The vast majority of aircraft NO_x emissions occur at these lower cruising altitudes of commercial subsonic aircraft (NASA 1996).⁵⁹

International marine bunkers comprise emissions from fuels burned by ocean-going ships of all flags that are engaged in international transport. Ocean-going ships are generally classified as cargo and passenger carrying, military (i.e., Navy), fishing, and miscellaneous support ships (e.g., tugboats). For the purpose of estimating greenhouse gas emissions, international bunker fuels are solely related to cargo and passenger carrying vessels, which is the largest of the four categories, and military vessels. Two main types of fuels are used on sea-going vessels: distillate diesel fuel and residual fuel oil. Carbon dioxide is the primary greenhouse gas emitted from marine shipping. In comparison to aviation, the atmospheric impacts of NO_x from shipping are relatively minor, as the emissions occur at ground level.

Overall, aggregate greenhouse gas emissions in 2003 from the combustion of international bunker fuels from both aviation and marine activities were 85.1 Tg CO₂ Eq., or 26 percent below emissions in 1990 (see Table 3-54). Although emissions from international flights departing from the United States have increased significantly (29 percent), emissions from international shipping voyages departing the United States have decreased by 63 percent since 1990. The majority of these emissions were in the form of CO₂; however, small amounts of CH₄ and N₂O were also emitted. Emissions of NO_x by aircraft during idle, take-off, landing and at cruising altitudes are of primary concern because of their effects on ground-level ozone formation (see Table 3-55).

Table 3-54: Emissions from International Bunker Fuels (Tg CO₂ Eq.)

Gas/Mode	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	113.5	109.9	114.6	105.3	101.4	97.9	89.5	84.2
Aviation	46.2	55.9	56.7	58.9	60.5	59.4	61.8	59.6
Marine	67.3	54.0	57.9	46.4	40.9	38.5	27.7	24.6
CH₄	0.2	0.1	0.2	0.1	0.1	0.1	0.1	0.1
Aviation	+	+	+	+	+	+	+	+
Marine	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
N₂O	1.0	1.0	1.0	0.9	0.9	0.9	0.8	0.8
Aviation	0.5	0.5	0.6	0.6	0.6	0.6	0.6	0.6
Marine	0.5	0.4	0.4	0.4	0.3	0.3	0.2	0.2
Total	114.6	111.0	115.7	106.4	102.4	98.9	90.4	85.1

+ Does not exceed 0.05 Tg CO₂ Eq.

⁵⁶ Naphtha-type jet fuel was used in the past by the military in turbojet and turboprop aircraft engines.

⁵⁷ Currently there are only around a dozen civilian supersonic aircraft in service around the world that fly at these altitudes, however.

⁵⁸ However, at this lower altitude, ozone does little to shield the earth from ultraviolet radiation.

⁵⁹ Cruise altitudes for civilian subsonic aircraft generally range from 8.2 to 12.5 km (27,000 to 41,000 feet).

Note: Totals may not sum due to independent rounding. Includes aircraft cruise altitude emissions.

Table 3-55: Emissions from International Bunker Fuels (Gg)

Gas/Mode	1990	1997	1998	1999	2000	2001	2002	2003
CO₂	113,503	109,858	114,557	105,294	101,404	97,865	89,489	84,193
Aviation	46,230	55,899	56,657	58,865	60,545	59,388	61,787	59,558
Marine	67,272	53,960	57,900	46,429	40,859	38,477	27,701	24,635
CH₄	8	7	7	6	6	5	4	4
Aviation	1	2	2	2	2	2	2	2
Marine	7	5	6	5	4	4	3	2
N₂O	3	3	3	3	3	3	3	2
Aviation	1	2	2	2	2	2	2	2
Marine	2	1	1	1	1	1	1	1
CO	115	124	127	124	124	120	118	113
Aviation	76	92	93	97	100	98	102	98
Marine	39	32	34	27	24	23	16	15
NO_x	1,985	1,668	1,778	1,478	1,334	1,266	988	900
Aviation	182	221	224	233	240	235	245	236
Marine	1,803	1,446	1,554	1,245	1,095	1,031	743	664
NMVOC	59	52	55	48	44	42	35	32
Aviation	11	14	14	15	15	15	15	15
Marine	48	38	41	33	29	27	20	18

Note: Totals may not sum due to independent rounding. Includes aircraft cruise altitude emissions.

Methodology

Emissions of CO₂ were estimated by applying of carbon content and fraction oxidized factors to fuel consumption activity data. This approach is analogous to that described under CO₂ from Fossil Fuel Combustion. Carbon content and fraction oxidized factors for jet fuel, distillate fuel oil, and residual fuel oil were taken directly from the EIA and are presented in Annex 2.1, Annex 2.2, and Annex 3.7. Heat content and density conversions were taken from EIA (2004) and USAF (1998). A complete description of the methodology and a listing of the various factors employed can be found in Annex 2.1. See Annex 3.7 for a specific discussion on the methodology used for estimating emissions from international bunker fuel use by the U.S. military.

Emission estimates for CH₄, N₂O, CO, NO_x, and NMVOCs were calculated by multiplying emission factors by measures of fuel consumption by fuel type and mode. Emission factors used in the calculations of CH₄, N₂O, CO, NO_x, and NMVOC emissions were obtained from the *Revised 1996 IPCC Guidelines* (IPCC/UNEP/OECD/IEA 1997). For aircraft emissions, the following values, in units of grams of pollutant per kilogram of fuel consumed (g/kg), were employed: 0.09 for CH₄, 0.1 for N₂O, 5.2 for CO, 12.5 for NO_x, and 0.78 for NMVOCs. For marine vessels consuming either distillate diesel or residual fuel oil the following values, in the same units, except where noted, were employed: 0.32 for CH₄, 0.08 for N₂O, 1.9 for CO, 87 for NO_x, and 0.052 g/MJ for NMVOCs. Activity data for aviation included solely jet fuel consumption statistics, while the marine mode included both distillate diesel and residual fuel oil.

Activity data on aircraft fuel consumption were collected from three government agencies. Jet fuel consumed by U.S. flag air carriers for international flight segments was supplied by the Bureau of Transportation Statistics (DOT 1991 through 2004). It was assumed that 50 percent of the fuel used by U.S. flagged carriers for international flights—both departing and arriving in the United States—was purchased domestically for flights departing from the United States. In other words, only one-half of the total annual fuel consumption estimate was used in the calculations. Data on jet fuel expenditures by foreign flagged carriers departing U.S. airports was taken from unpublished data collected by the Bureau of Economic Analysis (BEA) under the U.S. Department of Commerce (BEA 1991 through 2004). Approximate average fuel prices paid by air carriers for aircraft on international flights was taken from DOT (1991 through 2004) and used to convert the BEA expenditure data to gallons of fuel consumed. Data on U.S. Department of Defense (DoD) aviation bunker fuels and total jet fuel consumed by the

U.S. military was supplied by the Office of the Under Secretary of Defense (Installations and Environment), DoD. Estimates of the percentage of each Services' total operations that were international operations were developed by DoD. Military aviation bunkers included international operations, operations conducted from naval vessels at sea, and operations conducted from U.S. installations principally over international water in direct support of military operations at sea. Military aviation bunker fuel emissions were estimated using military fuel and operations data synthesized from unpublished data by the Defense Energy Support Center, under DoD's Defense Logistics Agency (DESC 2004). Together, the data allow the quantity of fuel used in military international operations to be estimated. Densities for each jet fuel type were obtained from a report from the U.S. Air Force (USAF 1998). Final jet fuel consumption estimates are presented in Table 3-56. See Annex 3.7 for additional discussion of military data.

Activity data on distillate diesel and residual fuel oil consumption by cargo or passenger carrying marine vessels departing from U.S. ports were taken from unpublished data collected by the Foreign Trade Division of the U.S. Department of Commerce's Bureau of the Census (DOC 1991 through 2004). Activity data on distillate diesel consumption by military vessels departing from U.S. ports were provided by DESC (2004). The total amount of fuel provided to naval vessels was reduced by 13 percent to account for fuel used while the vessels were not-underway (i.e., in port). Data on the percentage of steaming hours underway versus not-underway were provided by the U.S. Navy. These fuel consumption estimates are presented in Table 3-57.

Table 3-56: Aviation Jet Fuel Consumption for International Transport (Million Gallons)

Nationality	1990	1997	1998	1999	2000	2001	2002	2003
U.S. Carriers	1,954	2,457	2,462	2,625	2,737	2,619	2,495	2,418
Foreign Carriers	2,051	2,939	3,009	3,093	3,166	3,118	3,537	3,388
U.S. Military	862	496	502	488	480	524	482	473
Total	4,867	5,892	5,973	6,206	6,384	6,261	6,515	6,280

Note: Totals may not sum due to independent rounding.

Table 3-57: Marine Fuel Consumption for International Transport (Million Gallons)

Fuel Type	1990	1997	1998	1999	2000	2001	2002	2003
Residual Fuel Oil	4,781	3,843	3,974	3,272	2,967	2,846	1,937	1,597
Distillate Diesel Fuel & Other	617	421	627	308	290	204	158	137
U.S. Military Naval Fuels	522	484	518	511	329	318	348	459
Total	5,920	4,748	5,119	4,091	3,586	3,368	2,443	2,193

Note: Totals may not sum due to independent rounding.

Uncertainty

Emission estimates related to the consumption of international bunker fuels are subject to the same uncertainties as those from domestic aviation and marine mobile combustion emissions; however, additional uncertainties result from the difficulty in collecting accurate fuel consumption activity data for international transport activities separate from domestic transport activities.⁶⁰ For example, smaller aircraft on shorter routes often carry sufficient fuel to complete several flight segments without refueling in order to minimize time spent at the airport gate or take advantage of lower fuel prices at particular airports. This practice, called tankering, when done on international flights, complicates the use of fuel sales data for estimating bunker fuel emissions. Tankering is less common with the type of large, long-range aircraft that make many international flights from the United States, however. Similar practices occur in the marine shipping industry where fuel costs represent a significant portion of overall operating costs and fuel prices vary from port to port, leading to some tankering from ports with low fuel costs.

⁶⁰ See uncertainty discussions under Carbon Dioxide Emissions from Fossil Fuel Combustion.

Particularly for aviation, the DOT (1991 through 2004) international flight segment fuel data used for U.S. flagged carriers does not include smaller air carriers and unfortunately defines flights departing to Canada and some flights to Mexico as domestic instead of international. As for the BEA (1991 through 2004) data on foreign flagged carriers, there is some uncertainty as to the average fuel price, and to the completeness of the data. It was also not possible to determine what portion of fuel purchased by foreign carriers at U.S. airports was actually used on domestic flight segments; this error, however, is believed to be small.⁶¹

Uncertainties exist with regard to the total fuel used by military aircraft and ships, and in the activity data on military operations and training that were used to estimate percentages of total fuel use reported as bunker fuel emissions. Total aircraft and ship fuel use estimates were developed from DoD records, which document fuel sold to the Navy and Air Force from the Defense Logistics Agency. These data may slightly over or under estimate actual total fuel use in aircraft and ships because each Service may have procured fuel from, and/or may have sold to, traded with, and/or given fuel to other ships, aircraft, governments, or other entities. There are uncertainties in aircraft operations and training activity data. Estimates for the quantity of fuel actually used in Navy and Air Force flying activities reported as bunker fuel emissions had to be estimated based on a combination of available data and expert judgment. Estimates of marine bunker fuel emissions were based on Navy vessel steaming hour data, which reports fuel used while underway and fuel used while not underway. This approach does not capture some voyages that would be classified as domestic for a commercial vessel. Conversely, emissions from fuel used while not underway preceding an international voyage are reported as domestic rather than international as would be done for a commercial vessel. There is uncertainty associated with ground fuel estimates for 1997 through 2001. Small fuel quantities may have been used in vehicles or equipment other than that which was assumed for each fuel type.

There are also uncertainties in fuel end-uses by fuel-type, emissions factors, fuel densities, diesel fuel sulfur content, aircraft and vessel engine characteristics and fuel efficiencies, and the methodology used to back-calculate the data set to 1990 using the original set from 1995. The data were adjusted for trends in fuel use based on a closely correlating, but not matching, data set. All assumptions used to develop the estimate were based on process knowledge, Department and Component data, and expert judgments. The magnitude of the potential errors related to the various uncertainties has not been calculated, but is believed to be small. The uncertainties associated with future military bunker fuel emission estimates could be reduced through additional data collection.

Although aggregate fuel consumption data have been used to estimate emissions from aviation, the recommended method for estimating emissions of gases other than CO₂ in the *Revised 1996 IPCC Guidelines* is to use data by specific aircraft type (IPCC/UNEP/OECD/IEA 1997). The IPCC also recommends that cruise altitude emissions be estimated separately using fuel consumption data, while landing and take-off (LTO) cycle data be used to estimate near-ground level emissions of gases other than CO₂.⁶²

There is also concern as to the reliability of the existing DOC (1991 through 2004) data on marine vessel fuel consumption reported at U.S. customs stations due to the significant degree of inter-annual variation.

⁶¹ Although foreign flagged air carriers are prevented from providing domestic flight services in the United States, passengers may be collected from multiple airports before an aircraft actually departs on its international flight segment. Emissions from these earlier domestic flight segments should be classified as domestic, not international, according to the IPCC.

⁶² U.S. aviation emission estimates for CO, NO_x, and NMVOCs are reported by EPA's National Emission Inventory (NEI) Air Pollutant Emission Trends web site, and reported under the Mobile Combustion section. It should be noted that these estimates are based solely upon LTO cycles and consequently only capture near ground-level emissions, which are more relevant for air quality evaluations. These estimates also include both domestic and international flights. Therefore, estimates reported under the Mobile Combustion section overestimate IPCC-defined domestic CO, NO_x, and NMVOC emissions by including landing and take-off (LTO) cycles by aircraft on international flights, but underestimate because they do not include emissions from aircraft on domestic flight segments at cruising altitudes. The estimates in Mobile Combustion are also likely to include emissions from ocean-going vessels departing from U.S. ports on international voyages.

QA/QC and Verification

A source-specific QA/QC plan for international bunker fuels was developed and implemented. This effort included a Tier 1 analysis, as well as portions of a Tier 2 analysis. The Tier 2 procedures that were implemented involved checks specifically focusing on the activity data and emission factor sources and methodology used for estimating CO₂, CH₄, and N₂O from international bunker fuels in the United States. Emission totals for the different sectors and fuels were compared and trends were investigated. No corrective actions were necessary.

Recalculations Discussion

Historical activity data for aviation was slightly revised for both U.S. and foreign carriers. These changes were due to revisions to international fuel cost for foreign carriers and international jet fuel consumption for U.S. carriers, provided by DOT (1991 through 2004). These historical data changes resulted in minimal changes to the emission estimates for 1990 through 2002, which averaged to an annual increase in emissions from international bunker fuels of less than 0.1 Tg CO₂ Eq. (0.1 percent) in CO₂ emissions, annual increase of less than 0.1 Tg CO₂ Eq. (less than 0.1 percent) in CH₄ emissions, and annual increase of less than 0.1 Tg CO₂ Eq. (0.1 percent) in N₂O emissions.

3.12. Wood Biomass and Ethanol Consumption (IPCC Source Category 1A)

The combustion of biomass fuels—such as wood, charcoal, and wood waste—and biomass-based fuels—such as ethanol from corn and woody crops—generates CO₂. However, in the long run the CO₂ emitted from biomass consumption does not increase atmospheric CO₂ concentrations, assuming the biogenic carbon emitted is offset by the uptake of CO₂ resulting from the growth of new biomass. As a result, CO₂ emissions from biomass combustion have been estimated separately from fossil fuel-based emissions and are not included in the U.S. totals. Net carbon fluxes from changes in biogenic carbon reservoirs in wooded or crop lands are accounted for in the Land-Use Change and Forestry chapter.

In 2003, total CO₂ emissions from the burning of woody biomass in the industrial, residential, commercial, and electricity generation sectors were approximately 201.0 Tg CO₂ Eq. (201,042 Gg) (see Table 3-58 and Table 3-59). As the largest consumer of woody biomass, the industrial sector was responsible for 71 percent of the CO₂ emissions from this source. The residential sector was the second largest emitter, constituting 18 percent of the total, while the commercial and electricity generation sectors accounted for the remainder.

Table 3-58: CO₂ Emissions from Wood Consumption by End-Use Sector (Tg CO₂ Eq.)

End-Use Sector	1990	1997	1998	1999	2000	2001	2002	2003
Industrial	135.3	162.4	150.5	152.0	153.6	135.4	143.7	143.1
Residential	59.9	44.6	39.9	42.7	44.7	38.2	32.3	37.0
Commercial	4.0	5.0	5.0	5.4	5.5	4.2	4.3	4.4
Electricity Generation	13.3	14.1	14.1	14.2	13.9	13.0	15.5	16.6
Total	212.5	226.3	209.5	214.3	217.6	190.8	195.8	201.0

Note: Totals may not sum due to independent rounding.

Table 3-59: CO₂ Emissions from Wood Consumption by End-Use Sector (Gg)

End-Use Sector	1990	1996	1997	1998	1999	2000	2001	2002
Industrial	135,347	162,447	150,510	152,019	153,559	135,413	143,694	143,084
Residential	59,911	44,650	39,920	42,677	44,685	38,153	32,276	37,019
Commercial	4,037	5,042	4,963	5,394	5,481	4,175	4,319	4,369
Electricity Generation	13,252	14,126	14,097	14,233	13,851	13,034	15,487	16,570
Total	212,547	226,265	209,490	214,323	217,577	190,776	195,775	201,042

Note: Totals may not sum due to independent rounding.

Biomass-derived fuel consumption in the United States consisted primarily of ethanol use in the transportation sector. Ethanol is primarily produced from corn grown in the Midwest, and was used mostly in the Midwest and

South. Pure ethanol can be combusted, or it can be mixed with gasoline as a supplement or octane-enhancing agent. The most common mixture is a 90 percent gasoline, 10 percent ethanol blend known as gasohol. Ethanol and ethanol blends are often used to fuel public transport vehicles such as buses, or centrally fueled fleet vehicles. These fuels burn cleaner than gasoline (i.e., lower in NO_x and hydrocarbon emissions), and have been employed in urban areas with poor air quality. However, because ethanol is a hydrocarbon fuel, its combustion emits CO₂.

In 2003, the United States consumed an estimated 239 trillion Btus of ethanol, and as a result, produced approximately 15.8 Tg CO₂ Eq. (15,771 Gg) (see Table 3-60) of CO₂ emissions. Ethanol production and consumption has grown steadily every year since 1990, with the exception of 1996 due to short corn supplies and high prices in that year.

Table 3-60: CO₂ Emissions from Ethanol Consumption

Year	Tg CO ₂ Eq.	Gg
1990	4.2	4,155
1997	7.0	6,978
1998	7.7	7,711
1999	8.0	8,017
2000	9.2	9,188
2001	9.7	9,701
2002	11.5	11,473
2003	15.8	15,771

Methodology

Woody biomass emissions were estimated by taking U.S. consumption data (EIA 2004) (see Table 3-61), provided in energy units for the industrial, residential, commercial, and electric generation sectors, and applying two EIA gross heat contents (Lindstrom 2003). One heat content (16.953114 MMBtu/MT Wood & Wood Waste) was applied to the industrial sector's consumption, while the other heat content (15.432359 MMBtu/MT Wood & Wood Waste) was applied to the consumption data for the other sectors. An EIA emission factor of 0.434 MT C/MT Wood (Lindstrom 2003) was then applied to the resulting quantities of woody biomass to obtain CO₂ emissions estimates. It was assumed that the woody biomass contains black liquor and other wood wastes, has a moisture content of 12 percent, and is converted into carbon dioxide with 100 percent efficiency. The emissions from ethanol consumption were calculated by applying an EIA emission factor of 17.99 Tg C/QBtu (Lindstrom 2003) to U.S. ethanol consumption data that were provided in energy units (EIA 2004) (see Table 3-62).

Table 3-61: Woody Biomass Consumption by Sector (Trillion Btu)

Year	Industrial	Residential	Commercial	Electricity Generation
1990	1,442	581	39	129
1997	1,731	433	49	137
1998	1,603	387	48	137
1999	1,620	414	52	138
2000	1,636	433	53	134
2001	1,443	370	40	126
2002	1,531	313	42	150
2003	1,524	359	42	161

Table 3-62: Ethanol Consumption

Year	Trillion Btu
1990	63
1997	106
1998	117

1999	122
2000	139
2001	147
2002	174
2003	239

Uncertainty

It is assumed that the combustion efficiency for woody biomass is 100 percent, which is believed to be an overestimate of the efficiency of wood combustion processes in the United States. Decreasing the combustion efficiency would increase emission estimates. Additionally, the heat content applied to the consumption of woody biomass in the residential, commercial, and electric power sectors is unlikely to be a completely accurate representation of the heat content for all the different types of woody biomass consumed within these sectors. Emission estimates from ethanol production are more certain than estimates from woody biomass consumption due to better activity data collection methods and uniform combustion techniques.

Recalculations Discussion

The historical data for wood biomass consumption was adjusted slightly, which resulted in an average annual decrease in emissions from wood biomass and ethanol consumption of 0.3 Tg CO₂ Eq. (0.1 percent) from 1990 through 2002.

[BEGIN BOX]

Box 3-4: Formation of CO₂ through Atmospheric CH₄ Oxidation

Methane emitted to the atmosphere will eventually oxidize into CO₂, which remains in the atmosphere for up to 200 years. The global warming potential (GWP) of CH₄, however, does not account for the radiative forcing effects of the CO₂ formation that results from this CH₄ oxidation. The IPCC *Guidelines for Greenhouse Gas Inventories* (IPCC/UNEP/OECD/IEA 1997) do not explicitly recommend a procedure for accounting for oxidized CH₄, but some of the resulting CO₂ is, in practice, included in the inventory estimates because of the intentional “double-counting” structure for estimating CO₂ emissions from the combustion of fossil fuels. According to the IPCC Guidelines, countries should estimate emissions of CH₄, CO, and NMVOCs from fossil fuel combustion, but also assume that these compounds eventually oxidize to CO₂ in the atmosphere. This is accomplished by using CO₂ emission factors that do not factor out carbon in the fuel that is released as in the form of CH₄, CO, and NMVOC molecules. Therefore, the carbon in fossil fuel is intentionally double counted, as an atom in a CH₄ molecule and as an atom in a CO₂ molecule.⁶³ While this approach does account for the full radiative forcing effect of fossil fuel-related greenhouse gas emissions, the timing is not accurate because it may take up to 12 years for the CH₄ to oxidize and form CO₂.

There is no similar IPCC approach to account for the oxidation of CH₄ emitted from sources other than fossil fuel combustion (e.g., landfills, livestock, and coal mining). Methane from biological systems contains carbon that is part of a rapidly cycling biological system, and therefore any carbon created from oxidized CH₄ from these sources

⁶³ It is assumed that 100 percent of the CH₄ emissions from combustion sources are accounted for in the overall carbon emissions calculated as CO₂ for sources using emission factors and carbon mass balances. However, it may be the case for some types of combustion sources that the oxidation factors used for calculating CO₂ emissions do not accurately account for the full mass of carbon emitted in gaseous form (i.e., partially oxidized or still in hydrocarbon form).

is matched with carbon removed from the atmosphere by biological systems—likely during the same or subsequent year. Thus, there are no additional radiative forcing effects from the oxidation of CH₄ from biological systems. For example, the carbon content of CH₄ from enteric fermentation is derived from plant matter, which itself was created through the conversion of atmospheric CO₂ to organic compounds.

The remaining anthropogenic sources of CH₄ (e.g., fugitive emissions from coal mining and natural gas systems, industrial process emissions) do increase the long-term CO₂ burden in the atmosphere, and this effect is not captured in the inventory. The following tables provide estimates of the equivalent CO₂ production that results from the atmospheric oxidation of CH₄ from these remaining sources. The estimates for CH₄ emissions are gathered from the respective sections of this report, and are presented in Table 3-63. The CO₂ estimates are summarized in Table 3-64.

Table 3-63: CH₄ Emissions from Non-Combustion Fossil Sources (Gg)

Source	1990	1997	1998	1999	2000	2001	2002	2003
Coal Mining	3,900	2,983	2,989	2,805	2,677	2,647	2,497	2,561
Abandoned Coal Mines	288	385	341	349	369	331	303	306
Natural Gas Systems	6,112	6,363	6,276	6,066	6,289	6,277	6,221	5,998
Petroleum Systems	951	895	879	848	836	831	815	815
Petrochemical Production	56	78	80	81	80	68	72	72
Silicon Carbide Production	1	1	1	1	1	+	+	+
Iron and Steel Production	63	60	57	56	57	51	48	49
Total	11,371	10,765	10,622	10,205	10,308	10,206	9,956	9,801

Note: These emissions are accounted for under their respective source categories. Totals may not sum due to independent rounding.

Table 3-64: Formation of CO₂ through Atmospheric CH₄ Oxidation (Tg CO₂ Eq.)

Source	1990	1997	1998	1999	2000	2001	2002	2003
Coal Mining	10.7	8.2	8.2	7.7	7.4	7.3	6.9	7.0
Abandoned Coal Mines	0.8	1.1	0.9	1.0	1.0	0.9	0.8	0.8
Natural Gas Systems	16.8	17.5	17.3	16.7	17.3	17.3	17.1	16.5
Petroleum Systems	2.6	2.5	2.4	2.3	2.3	2.3	2.2	2.2
Petrochemical Production	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2
Silicon Carbide Production	+	+	+	+	+	+	+	+
Iron and Steel Production	0.2	0.2	0.2	0.2	0.2	0.1	0.1	0.1
Total	31.3	29.6	29.2	28.1	28.3	28.1	27.4	27.0

Note: Totals may not sum due to independent rounding.

+ Does not exceed 0.05 Tg CO₂ Eq.

The estimates of CO₂ formation are calculated by applying a factor of 44/16, which is the ratio of molecular weight of CO₂ to the molecular weight of CH₄. For the purposes of the calculation, it is assumed that CH₄ is oxidized to CO₂ in the same year that it is emitted. As discussed above, this is a simplification, because the average atmospheric lifetime of CH₄ is approximately 12 years.

Carbon dioxide formation can also result from the oxidation of CO and NMVOCs. However, the resulting increase of CO₂ in the atmosphere is explicitly included in the mass balance used in calculating the storage and emissions from non-energy uses of fossil fuels, with the carbon components of CO and NMVOC counted as CO₂ emissions in the mass balance.⁶⁴

[END BOX]

⁶⁴ See Annex 2.3 for a more detailed discussion on accounting for indirect emissions from CO and NMVOCs.

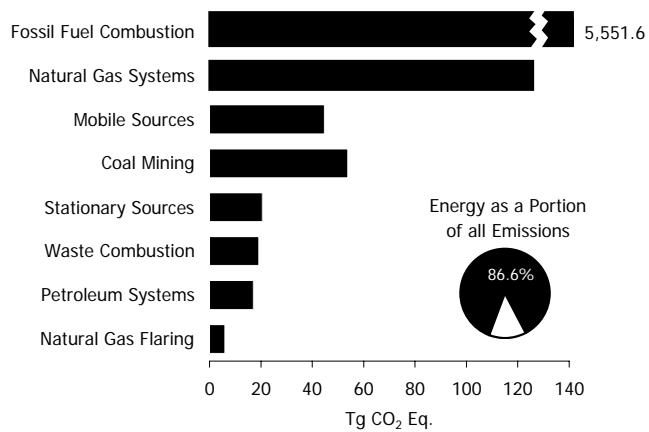
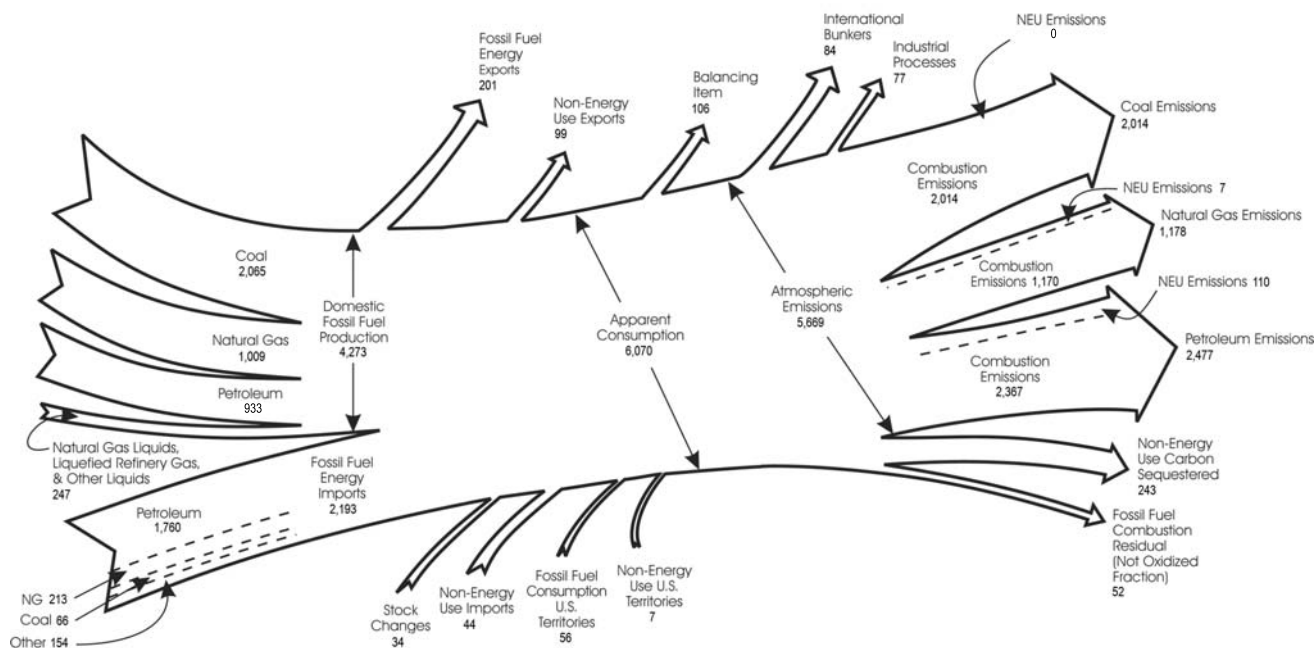


Figure 3-1: 2003 Energy Sector Greenhouse Gas Sources

Figure 3-2
2003 U.S. Fossil Carbon Flows (Tg CO₂ Eq.)



Note: Totals may not sum due to independent rounding.

The "Balancing Item" above accounts for statistical imbalances and unknowns in the reported data sets combined here.

NEU = Non-Energy Use
 NG = Natural Gas

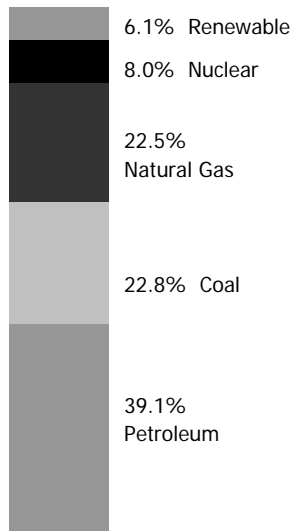


Figure 3-3: 2003 U.S. Energy Consumption by Energy Source

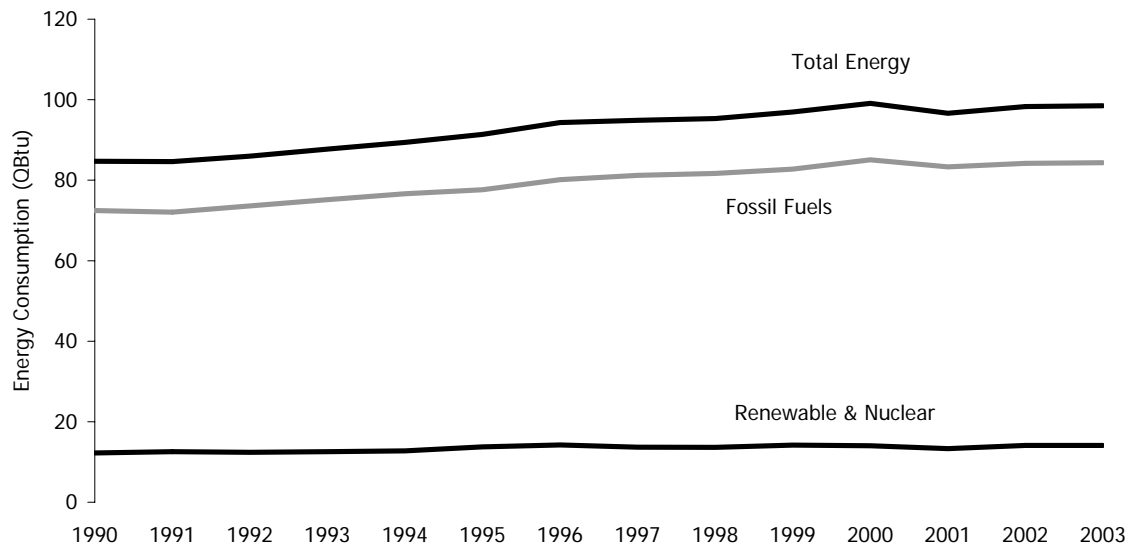


Figure 3-4: U.S. Energy Consumption (Quadrillion Btu)

Note: Expressed as gross calorific values.

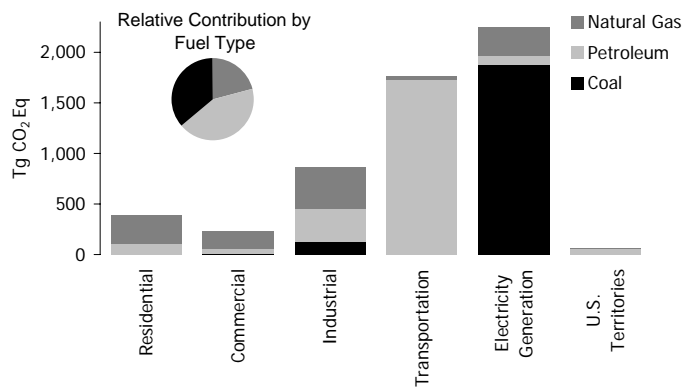


Figure 3-5: 2003 CO₂ Emissions from Fossil Fuel Combustion by Sector and Fuel Type

Note: The electricity generation sector also includes emissions of less than 0.01 Tg CO₂ Eq. from geothermal-based electricity generation

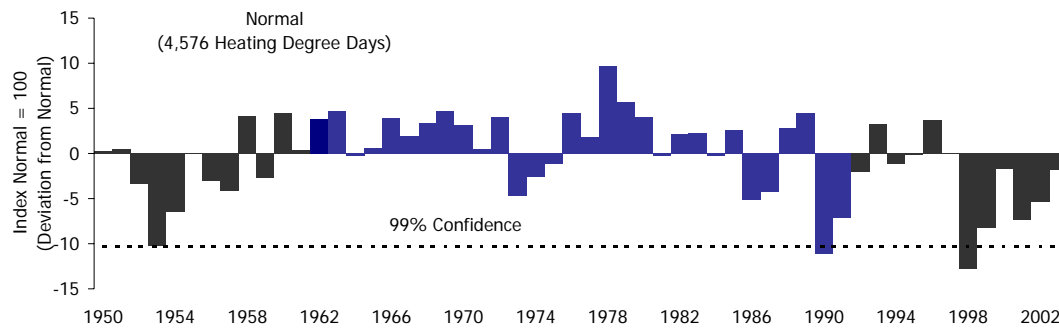


Figure 3-6. Annual Deviations from Normal Heating Degree Days for the United States (1949-2003)
 Note: Climatological normal data are highlighted.
 Statistical confidence interval for "normal" climatology period of 1961 through 1990.

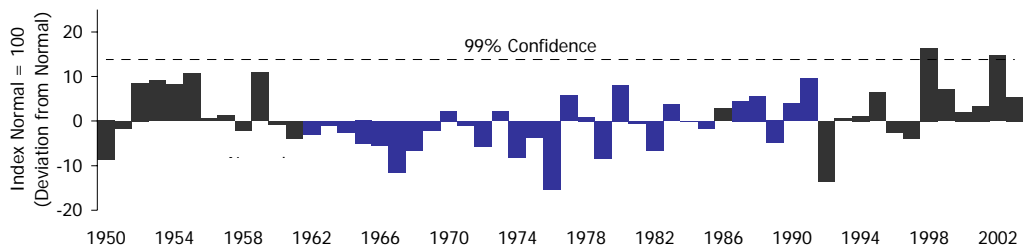


Figure 3-7: Annual Deviations from Normal Cooling Degree Days for the United States (1949-2003)
 Note: Climatological normal data are highlighted.
 Statistical confidence interval for "normal" climatology period of 1961 through 1990.

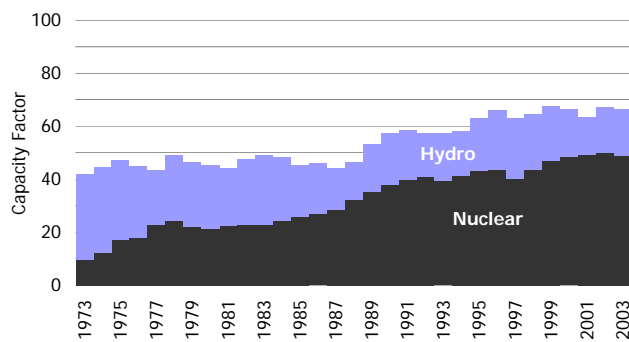


Figure 3-8: Aggregate Nuclear and Hydroelectric Power Plant Capacity Factors in the United States (1973-2003)

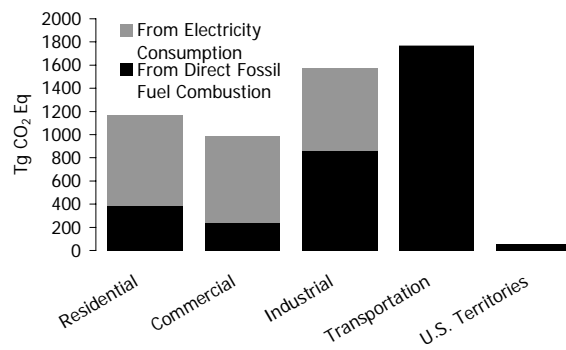


Figure 3-9: 2003 End-Use Sector Emissions of CO₂ from Fossil Fuel Combustion

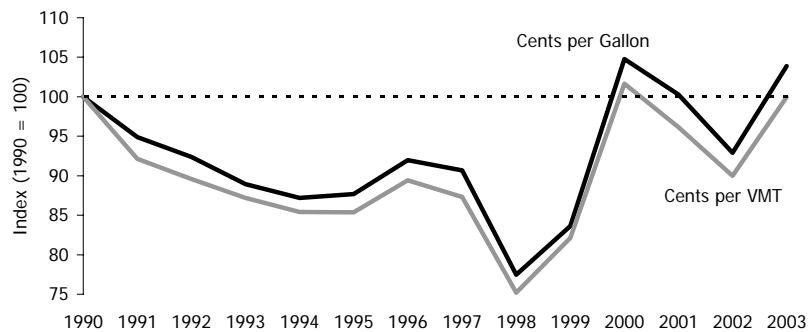


Figure 3-10: Motor Gasoline Retail Prices (Real)

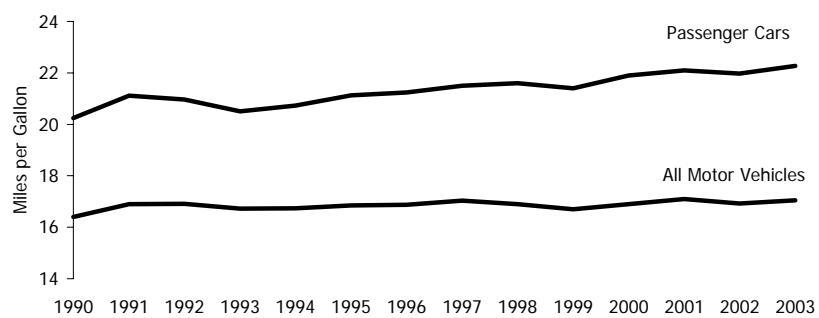


Figure 3-11: Motor Vehicle Fuel Efficiency

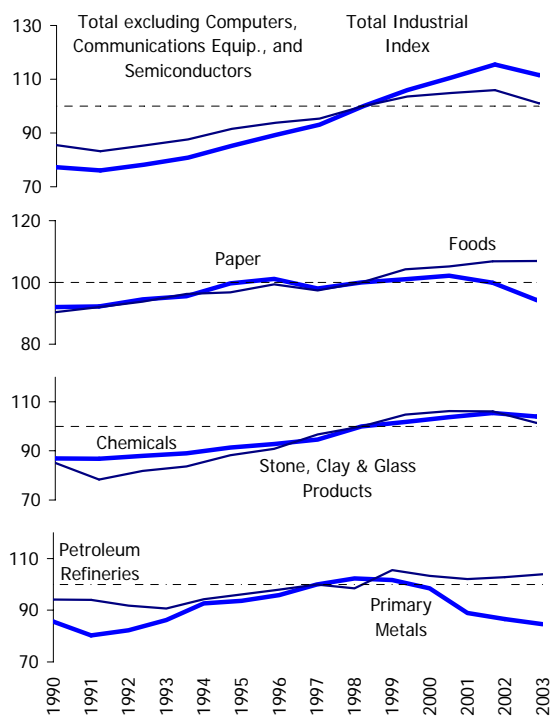


Figure 3-12: Industrial Production Indexes (Index 1997=100)

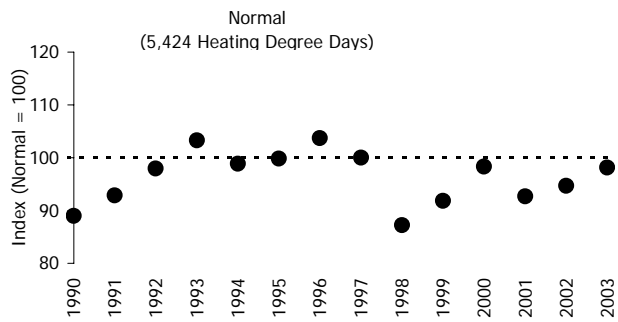


Figure 3-13: Heating Degree Days
Note: Excludes Alaska and Hawaii

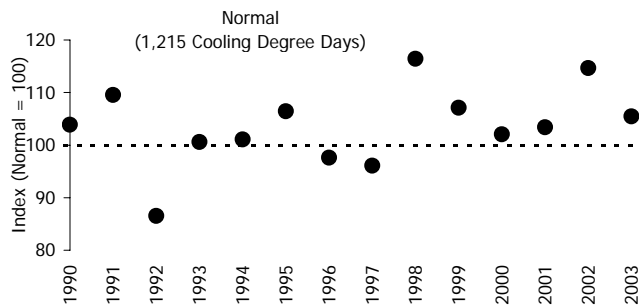


Figure 3-14: Cooling Degree Days
Note: Excludes Alaska and Hawaii

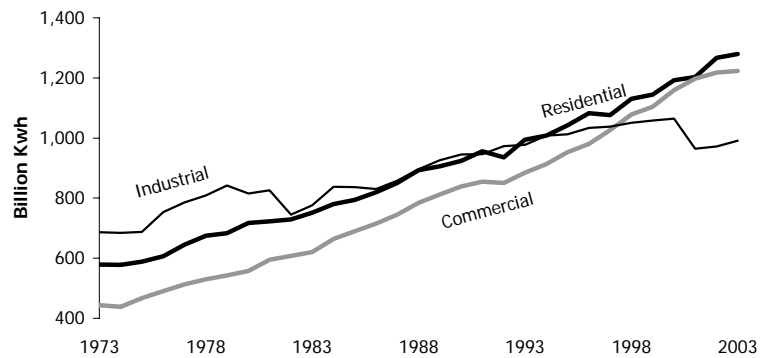


Figure 3-15: Electric Generation Retail Sales by End-Use Sector
Note: The transportation end-use sector consumes minor quantities of electricity.

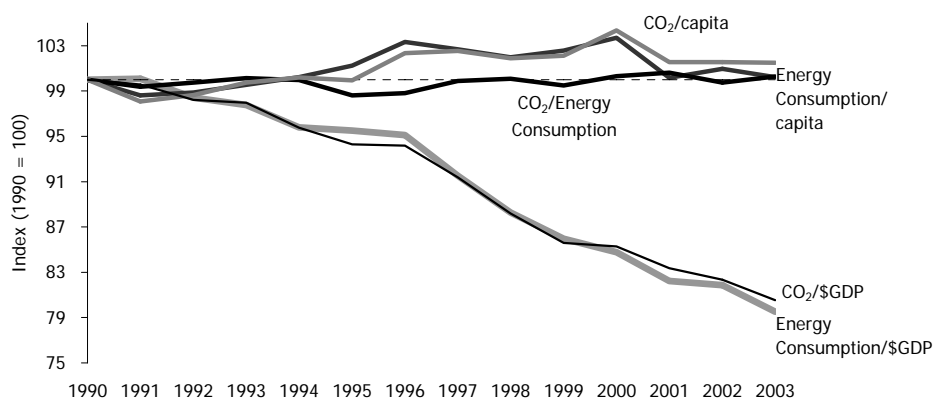


Figure 3-16: U.S. Energy Consumption and Energy-Related CO₂ Emissions Per Capita and Per Dollar GDP

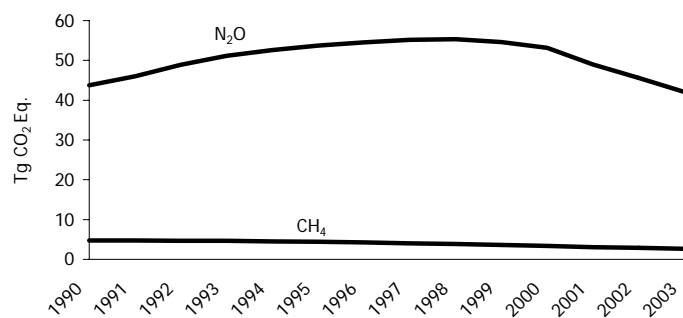


Figure 3-17: Mobile Source CH₄ and N₂O Emissions